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This disclosure relates to the forming of mineralized coatings on metal surfaces and to methods of forming such coatings. The coating include a wide range of compounds and normally at least a portion of the coating corresponds to an amorphous phase. The coating and method are particularly useful in providing a corrosion resistant coating or film upon a metallic surface. This aspect of the disclosure involves the formation of a coated retentate "hydrocolloid" of selected components.

the metal substrate, or to the metal substrate.

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CORROSION RESISTANT COATINGS CONTAINING AN AMORPHOUS PHASE

CROSS-REFERENCE TO RELATED APPLICATIONS

This application is a continuation in part of United States Patent Application Serial Nos. 08/850,586 (Attorney Docket No. EL001RH-7) and 08/850,523 (Attorney Docket No. EL001RH-6), filed on May 2, 1997; Serial Nos. 08/791,336 and 08/791,336, filed on January 31, 1997 (Attorney Docket No. EL001RH-4 and EL001RH-5); 08/634,215 (Attorney Docket No. EL001RH-3), entitled "Corrosion Resistant Buffer System for Metal Products" and filed on April 18, 1996 in the names of Robert L. Heimann, et al., which is a continuation in part of US Patent Application Serial No. 08/476,271 (Attorney Docket EL001RH-2) filed on June 7, 1995, which in turn is a continuation-in-part of US Patent Application Serial No. 08/327,438 (Attorney Docket EL001RH-1) filed on October 21, 1994, now allowed. The disclosure of each of the previously identified patent applications is hereby incorporated by reference.

5 vehicles, automobiles, aircraft, among a wide range of metallic components.

Current industry standards for corrosion prevention center around the use of barrier coatings, sacrificial coatings, alloys containing heavy metals such as chromium, nickel, lead, cadmium, copper, mercury, barium, among other heavy metals. The introduction of these materials into the environment, however, can lead to serious health consequences as well as substantial costs to contain or separate the materials or clean up environmental contamination. Damage associated with corrosion, accordingly, is a continuing problem and better systems for preventing corrosion are still needed.

A more detailed discussion of mineral containing materials can be found in Manual of Mineralogy, by Dana, 21st edition, American Society of Metals, vol. 13, *Composition in Structures, Reaction Sequence in Atmospheric Corrosion of Zinc ASTM STP 1239*, by W. W. Kirk et al., and Physics and Chemistry of Mineral Surfaces, by Bradley (1996); the disclosure of each of the aforementioned references is hereby incorporated by reference.

Conventional practices for improving protecting metal containing surfaces and imparting improved surface characteristics to metals rely upon compositions and methods which are undesirable as being costly or environmentally unsound.

FIELD OF THE INVENTION

The instant invention relates to the forming coatings on metal containing surfaces and to methods of forming such coatings on a suitable substrate. The coating can include a wide range of compounds and normally at least a portion of the coating corresponds to an amorphous phase. The inventive coating and method are particularly useful in providing a corrosion resistant coating or film upon a metallic surface. This aspect of the invention involves the formation of a corrosion resistant "mineralized" layer of tailored composition upon a metal containing surfaces.

BACKGROUND OF THE INVENTION

The corrosion of steel and other metal containing products continues to be a serious technical problem which has profound effects on the economy. Corrosion causes loss of natural resources, and deteriorates key infrastructure such as roads and buildings. It also causes premature replacement of equipment and parts in industrial facilities, boats and other marine

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coatings, sacrificial coatings, alloys containing heavy metals such as chromium, nickel, lead, cadmium, copper, mercury, barium, among other heavy metals. The introduction of these materials into the environment, however, can lead to serious health consequences as well as substantial costs to contain or separate the materials or clean up environmental contamination. Damage associated with corrosion, accordingly, is a continuing problem and better systems for preventing corrosion are still needed.

A more detailed discussion of mineral containing materials can be found in Manual of Mineralogy, by Dana, 21st edition, American Society of Metals, vol. 13, *Composition in Structures, Reaction Sequence in Atmospheric Corrosion of Zinc ASTM STP 1239*, by W. W. Kirk et al., and Physics and Chemistry of Mineral Surfaces, by Bradley (1996); the disclosure of each of the aforementioned references is hereby incorporated by reference.

Conventional practices for improving protecting metal containing surfaces and imparting improved surface characteristics to metals rely upon compositions and methods which are undesirable as being costly or environmentally unsound.

SUMMARY OF THE INVENTION

The instant invention solves problems associated with conventional practices by providing an improved method and a composition for improving the surface characteristics of a metal containing surface. While the inventive composition is normally compatible with conventional compositions and methods, the inventive composition can obviate the need to employ heavy metals such as chrome and environmentally undesirable solvents.

The present invention in a broad aspect relates to compositions and methods for improving or modifying the surface characteristics of a metal containing surface. In one aspect, the invention involves methods for forming a "mineralized" layer upon the surface of a substrate. One method of forming the mineralized layer comprises delivering precursors of the mineralized layer to the surface of the metal surface via a carrier. The carrier can be a wide range of known compositions such as a film forming composition, lubricants, gel, sealant, adhesive, paint, solvent and water-borne resins, among other conventional compositions for forming coatings or films upon metals. If

5 desired, the carrier can function as a reservoir of precursor materials thereby permitting additional formation of the mineralized layer, e.g., when in the presence of a reservoir a breach in the mineralized layer can be overcome by secondary mineral formation from mineral precursors in the reservoir - a so-called self healing effect. If desired, the carrier can also function as a reservoir of buffer materials, e.g., materials that passivate the pH of the metal surface, which can protect the metal surface by providing an environment in which the metal is resistant to chemical attack.

10 Depending upon the utility of the carrier, the carrier can be removed or remain permanently in contact with the mineralized surface (and at least a portion of the metal surface).

The instant invention provides an improved surface on articles by tailoring the surface chemistry and effecting a new mineralized surface through chemical reaction and interaction.

15 The mineralized surface is formed when precursors are delivered to the surface of a metal or metal coated articles or substrates. In some cases, the carrier includes materials which can function to buffer the surface, as a precursor of the mineralized layer, alter pH, activate the surface by changing the surface chemical species, or all of these functions. In some cases, the surface mineralization is enhanced by a pretreatment step prior to application of the precursor-containing carrier. After providing a proper environment, precursors can interact thereby in situ forming the mineralized layer upon at least a portion of the metal surface. Depending upon the surface environment, the metal or metal coated substrate can contribute donor ions to react and/or interact with delivered precursors thereby forming a relatively thin mineralized layer that is effective in altering and preferably enhance the characteristics of the entire article, e.g., by altering and preferably enhancing the surface characteristics of the article. Consequently, the instant invention permits tailoring a metal containing surface to possess improved corrosion, coating adhesion, chemical resistance, thermal resistance, mechanical abrasion, acid rain resistance, UV resistance, resistance to effects from atomic oxygen and vacuum UV, engineered electrical resistance, among other improved properties. As will be described below in greater detail, at least a portion of the mineralized coating or layer normally corresponds to a novel amorphous phase.

5 coating or film upon at least a portion of a metal containing surface. By "mineralized" it is meant a composition containing at least one member selected from the group of oxygenated cations and anions wherein at least a portion of the mineral corresponds to an amorphous phase or matrix that embeds or surrounds an inorganic complex oxide crystal. Normally, the amorphous phase is the predominant phase component of the mineralized layer and, in some cases, substantially transparent to visible light. This type of predominantly amorphous structure is characterized by continuous random network (CRN). In general, the mineral layer has a network structure in which metal atoms are bound to oxygen atoms by predominantly covalent bonds. The CRN or network can be modified by the introduction of metal oxides which are, in many embodiments of the invention, contributed by the substrate. By "metal containing surface", "substrate", or "surface", it is meant to refer to a metallic article and any metal containing surface as well as any substrate at least partially coated with a metal layer, film or foil including a non-metallic article having a metal layer. A wide variety of substances can be employed as precursors of the mineralized layer, such as one or more cations of the metals of Groups I, II and III, and the transition metals, of the Periodic Chart of the Elements. Additionally, one or more of the anions selected from the group consisting of water soluble salts and/or oxides of tungsten, molybdenum, chromium, titanium, zircon, vanadium, phosphorus, aluminum, iron, boron, bismuth, gallium, tellurium, germanium, antimony, niobium (also known as columbium), magnesium and manganese, mixtures thereof, among others. Particularly desirable results can be obtained by using salts and oxides of silicon, aluminum and iron.

10 At least a portion of the resulting mineralized layer having oxide network attributes can be characterized by the following formula:

$$A_x B_y O_z \cdot nH_2O$$

where A is termed a modifier cation and may be one or more ions selected from Group I, II and/or III metals, and B is a network forming cation, such as silicon, aluminum, iron or magnesium. The values of x and y can be any number except zero but x and y cannot all concurrently be zero and z cannot be zero. "z" can have any value ranging from y to 4y. "n" is water of hydration and has a value of from about 0 to about 10. The relationships of x, y, and z follow rules by Zachariasen in the Journal of the American Chemical Society, Volume 54, page 3841 (1932); hereby incorporated by reference:

DETAILED DESCRIPTION

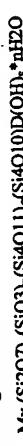
The instant invention relates to compositions and methods for forming a mineralized

5 1). A high proportion of (network-forming cations are surrounded by oxygen tetrahedra or triangles.

2). The oxygen polyhedra only share corners with each other.

3) Some oxygen atoms are linked to only two cations, and do not form additional bonds with any other cations.

10 In one aspect of the invention, the complex inorganic crystals that are surrounded by or incorporated within the amorphous matrix to form the mineral layer can also be characterized by the following formula:



wherein M_x is one or more metals supplied by the previously described substrate and x ranges from 1 to 5; $A + B + C + D = 1$ wherein A, B, C and D can each individually equal zero but cannot simultaneously be zero; n is the water of hydration and ranges from 0 to 10 and typically ranges from 0 to 6; and t, u, v and w are integers that ranges from about 0 to about 4.

In another aspect of the invention, at least a portion of the crystalline component of the mineral layer that is surrounded or incorporated within the amorphous phase comprises:



where M_x, M'_y and M''_z are ions of Group I, II and/or III metals, and A and A' are the previously defined anions and where x, y , and z can be any number including zero but x, y and z cannot all concurrently be zero. Analogously, t, u, v, w and x can each be any number including zero but cannot all concurrently be zero. t, u, v, w and x can each be any number including zero but about 10. At least one of M, M' and M'' is a metal supplied from the substrate in contact with the mineralized layer, and normally up to two of M, M' or M'' corresponds to an alkali or alkaline earth metal, e.g. calcium, potassium, sodium and mixtures thereof. For example, when the metal substrate comprises zinc and a precursor comprises sodium silicate the crystalline component, which is embedded within the amorphous matrix to form the mineralized layer, comprises



In a further aspect of the invention, at least a portion of the crystalline component of the mineral layer that is surrounded or incorporated within the amorphous phase comprises:



5 where M, M' , and M'' are ions of Group I, II and/or III metals, and A and A' are the previously defined anions and where v, w, x, y , and z can be any number including zero but x, y and z cannot all concurrently be zero. Analogously, A, B, C and D can each be any number including zero but cannot all concurrently be zero. $“n”$ is the water of hydration and normally ranges from about 0 to about 10; and typically, ranges from about 0 to 6. $“S”$ is an integer that ranges from about 0 to about 4. At least one of M, M' and M'' is a metal supplied from the substrate in contact with the mineralized layer, and normally up to two of M, M' or M'' corresponds to an alkali or alkaline earth metal, e.g. calcium, potassium, sodium and mixtures thereof. Without wishing to be bound by any theory or explanation, it is believed that the presence of alkali cations, e.g. M'' , can influence the presence of other metal ions, e.g., M' supplied from the metal substrate, by an exchange or a replacement mechanism. For example, when the metal substrate comprises zinc and a precursor comprises sodium silicate the crystalline component, which is embedded within the amorphous matrix to form the mineralized layer, comprises $Zn_xNa_yMg_z(Si_2O_7)_k(D(OH)_3 \cdot nH_2O.$

10 The mineralized layer is formed from precursors. By “precursors” it is meant any combination of materials which interact with the metal surface or substrate to form the mineralized layer as well as intermediate products that interact further to form the mineralized layer. Examples of precursors include buffers such as silicate buffers and carbonate buffers including sodium hydroxide; alkali silicates such as at least one of sodium, calcium and potassium silicate; silica; cations supplied or delivered to the surface such as at least one of zinc, molybdenum; ions supplied or delivered to the surface such as at least one of oxygen, sulfur or chlorine from the environment surrounding the precursors or surface; compounds which decompose or react to form a precursor or intermediate thereof; mixtures thereof, among others. Depending upon the precursors, amount thereof and contact conditions, a silica containing layer may form upon the mineralized layer.

15 The mineralized layer can be tailored by adding one or more dopants to the precursor. Examples of suitable dopants comprise at least one member from the group consisting of anions selected from the group consisting of water soluble salts and/or oxides of tungsten, molybdenum, chromium, titanium, zircon, vanadium, phosphorus, aluminum, iron, boron, bismuth, gallium, tellurium, germanium, antimony, niobium (also known as columbium), magnesium and manganese, mixtures thereof, among others, and more especially, salts and oxides of aluminum and iron. Desirable results can be obtained by adding one or more dopants to a sodium silicate

5 precursor. The precursors of the mineralized layer are added to any suitable carrier. Examples of suitable carriers include hydrocarbons such as at least one member selected from the group consisting of animal, vegetable, petroleum derived and synthetic oils such as polyalphaolefin (PAO), silicone oil, phosphate esters, fluorinated oils such as KRYTOX (supplied by the DuPont Company). Further examples of suitable carriers comprise at least one member selected from the group consisting of thermoplastic, thermosetting, cross-linked system, mixtures thereof, among others. Specific examples of such carriers include epoxies, acrylics, polyurethanes, silicones, polyesters, allyls, vinyls, phenolics, fluoropolymers, latexes, mixtures thereof, among others. Depending upon the process conditions, the precursor carrier may be selected from alkylated aromatics, phosphate esters, perfluoralkylpolyethers, polyesters, olefins, chlorotrifluoroethylene, silahydrocarbons, phosphazenes, diarylcarbonates, oligomers, polybutenes, and polyphenyl esters, as well as unsaturated polyglycols, silicones, silicate esters, cycloaliphatic hydrocarbons, and dibasic acid esters, e.g., when applying a precursor carrier to an iron containing surface a polyalphaolefin base oil having a kinematic viscosity in the range of about 30 - 1,400 centistokes at 40°C can be employed. Other properties to consider when choosing an appropriate polyalphaolefin base oil are molecular weight, molecular branching, thermal stability, and hydrophobicity, depending on the application. The polyalphaolefin base oil can be thickened to a gel with thickeners known to the art of grease manufacturers such as polytetrafluoroethylene or silica. Buffer materials are also suitable as thickeners as long as they are compatible with the base oil. 10 Generally, low molecular weight, synthetic, hydrocarbon oils provide greater ease in designing and manufacturing a gel with particular desired characteristics but are more costly than less refined, high molecular weight, petroleum hydrocarbon oils. Less refined hydrocarbons may also have the disadvantage of containing sulfide compounds which can feed sulfate reducing bacteria and, in turn, tend to corrode metals such as steel, iron, and iron alloys.

15 The carrier film or layer can have a thickness of about 1 to at least about 50 mils, and typically has a thickness of about 1 to about 1.5 mil, e.g., about 0.2 to at least 0.4 mil. Normally, the carrier is semipermeable thereby permitting anions from the surrounding environment to contact precursors to the mineralized product. By "semipermeable" it is meant to refer to a microporous structure, either natural or synthetic allowing passage of ions, water and other solvent

5 molecules, and very small other molecules. The resin can be essentially insoluble in water and resistant to macro-penetration by flowing water. The resin layers, however, are normally permeable to water molecules and inorganic ions such as metal ions and silicate ions.

10 The amount of mineralized layer precursor present in the carrier typically comprises about 1 to about 60 wt.% of the carrier, e.g., normally about 5 to about 10 wt.% depending upon the carrier.

15 The mineralized layer precursors can be combined with the carrier in any suitable conventional manner known in this art.

20 The mineralized layer precursors can include or be employed along with one or more additives such as the pH buffers such as those listed below in Tables A and B, mixtures thereof, among others. In some cases, a buffer also functions as a precursor, e.g., sodium silicate. The amount of these additives typically ranges from about 1 to about 60 wt.% of the carrier, e.g., normally about 5 to about 10 wt.%. These additives can be added to the carrier in order to tailor the characteristics of the mineralized layer, the carrier itself, upon a pre-treated surface, among other characteristics. By adding suitable mineralized layer precursors, carrier, additives, among other materials, the surface of the metal containing layer can be tailored by forming a mineralized layer to possess improved corrosion resistance, adhesion, among other characteristics.

Table A - Examples of Buffering Compounds

Chemical Name	Formula
Boric Acid	H_3BO_3
Citric Acid	$H_3C_6H_5O_7 \cdot H_2O$
Sodium Hydroxide	$NaOH$
Trisodium Phosphate Dodecylate	$Na_3PO_4 \cdot 12H_2O$
Potassium Silicate	SiO_4K_2O 1.6-2.5 wt. Ratio
Sodium Silicate	SiO_4Na_2O 2.0-3.22 wt. Ratio

Chemical Name	Formula
Potassium Hydrogen Phthalate	KHC ₈ O ₄ H ₄
Potassium Dihydrogen Phosphate	KH ₂ PO ₄
Borax	Na ₂ B ₄ O ₇
Sodium Hydrogen Carbonate	NaHCO ₃
Disodium Phosphate Dodecahydrate	Na ₂ HPO ₄ · 12H ₂ O
Sodium Acetate	NaOOCCH ₃
Disodium Phosphate	Na ₂ HPO ₄

Table B - Examples of Weight Ratios of Buffering Components for Various pH Values

Desired pH	Weight	Chemical	Weight	Chemical	Weight	Chemical	Weight	Chemical	Weight	Chemical
4.5	1.00	Sodium Hydroxide	29.30	Potassium Hydrogen Phthalate						
5.0	1.00	Sodium Hydroxide	11.30	Potassium Hydrogen Phthalate						
5.5	1.00	Sodium Hydroxide	6.97	Potassium Hydrogen Phthalate						
6.0	1.00	Sodium Hydroxide	30.40	Potassium Dihydrogen Phosphate						
6.5	1.00	Sodium Hydroxide	12.20	Potassium Dihydrogen Phosphate						
7.0	1.00	Sodium Hydroxide	5.84	Potassium Dihydrogen Phosphate						
7.5	1.00	Sodium Hydroxide	4.14	Potassium Dihydrogen Phosphate						
8.0	1.00	Sodium Hydroxide	3.64	Potassium Dihydrogen Phosphate						
8.5	1.00	Boric Acid	0.84	Citric Acid	4.80	Trisodium Phosphate (12 H ₂ O)				
9.0	1.00	Boric Acid	0.84	Citric Acid	5.82	Trisodium Phosphate (12 H ₂ O)				
9.5	1.00	Sodium Hydroxide	13.55	Borax						

Desired pH	Weight	Chemical	Weight	Chemical	Weight	Chemical
		Hydroxide				
10.0	1.00	Sodium Hydroxide	6.52	Borax		
10.5	1.00	Sodium Hydroxide	5.25	Borax		
11.0	1.00	Sodium Hydroxide	2.31	Sodium Hydrogen Carbonate		
11.5	1.00	Sodium Hydroxide	8.00	Disodium Acid Phosphate (12H ₂ O)		
12.0	1.00	Sodium Hydroxide	1.30	Disodium Acid Phosphate (12H ₂ O)		
12.5	1.00	Sodium Hydroxide	15.00	Disodium Acid Phosphate		
13.0	1.00	Sodium Hydroxide	1.00	Sodium Acetate		

5 results can be obtained when forming a mineralized layer upon a zinc containing surface.

In an aspect of the invention, the metal containing surface comprises zinc which is contacted with a carrier including sodium silicate. The carrier can comprise PAO or polyurethane. The amount of sodium silicate within a PAO carrier typically ranges from about 1 to about 30 wt. %, e.g., about 5 to about 10 wt. %, whereas for a carrier comprising polyurethane the amount of sodium silicate typically ranges from about 1 to about 15 wt. %, e.g., about 5 to about 10 with especially desirable results being obtained at about 6.5 wt. % sodium silicate. When the previously desired carrier is employed, desirable results can be obtained by applying one or more topcoats to the carrier, e.g., polyurethane, polytetrafluoroethylene, mixtures thereof, among others. The topcoat can function as a physical barrier to the surrounding environment thereby providing further corrosion resistance. The thickness of the carrier layer including any topcoat normally ranges from about 0.75 to about at least about 1.5 mils. The corrosion resistance can be further enhanced by heat treating the coated metal surface. That is, after applying a silicate containing carrier to a zinc containing surface and allowing the carrier to incubate, the coated surface is heated (in any suitable atmosphere such as air), to a temperature of about 125 to about 175 C. Consequently, the instant invention permits tailoring the carrier, mineralization precursors, topcoat as well as any heat treatment to obtain a predetermined corrosion resistance, e.g., in the case of a zinc containing surface the ASTM B117 resistance can range from 100 to 3,000 hours.

In an aspect of the invention, the metal containing surface comprises zinc which is contacted with a carrier including sodium silicate. The carrier can comprise PAO or polyurethane. The amount of sodium silicate within a PAO carrier typically ranges from about 1 to about 30 wt. %, e.g., about 5 to about 10 wt. %, whereas for a carrier comprising polyurethane the amount of sodium silicate typically ranges from about 15 wt. %, e.g., about 5 to about 10 with especially desirable results being obtained at about 6.5 wt. % sodium silicate. When the previously desired carrier is employed, desirable results can be obtained by applying one or more topcoats to the carrier, e.g., polyurethane, polytetrafluoroethylene, mixtures thereof, among others. The topcoat can function as a physical barrier to the surrounding environment thereby providing further corrosion resistance. The thickness of the carrier layer including any topcoat normally ranges from about 0.75 to about at least about 1.5 mils. The corrosion resistance can be further enhanced by heat treating the coated metal surface. That is, after applying a silicate containing carrier to a zinc

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The aforementioned carrier can be applied to a metal containing surface by using any expedient method. Depending upon the desired results, the metal containing surface can be applied or repainted as appropriate. Examples of suitable methods for applying the tailored carrier comprise at least one of priming, spraying, dipping, roweling, among other conventional methods. By employing a suitable tailored carrier, the instant invention can form a mineralized layer 10 to protect a metal containing surface having at least one member from the group of magnesium, aluminum, vanadium, calcium, beryllium, manganese, cobalt, nickel, copper, lead, copper, brass, bronze, zirconium, thallium, chromium, zinc, alloys thereof, among others. Particularly, desirable

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5 comming surface and allowing the carrier to incubate, the coated surface is heated (in any suitable atmosphere such as air), to a temperature of about 125 to about 175 C. Consequently, the instant invention permits tailoring the carrier, mineralization precursors, topcoat as well as any heat treatment to obtain a predetermined corrosion resistance, e.g., in the case of a zinc containing surface the ASTM B117 resistance can range from 100 to 3,000 hours.

10 Without wishing to be bound by any theory or explanation, it is believed that the mineralized layer is formed under a variety of chemical and physical forces including 1) transporting ions through the carrier via osmotic pressure and diffusion thereby providing ions to the metal surface, 2) oxygen deprived environment, 3) buffering to provide a predetermined alkaline pH environment that is effective for formation of the mineralized layer upon a given metal surface, e.g. in the case of a zinc containing surface about 9.5 to at least about 10.5 pH, 4) heterogeneous process using any available ions, 5) water present at the surface, in the carrier or as a reaction product can be removed via heat, vacuum or solvent extraction, 6) using a reservoir adjacent to the metal surface that can control that ion transport rate as well as the rate of water (and moieties) passing through the reservoir and serve to provide, as needed, a continuous supply mineralized layer precursors, among other forces.

15 The process for forming the inventive mineralized layer can be initiated by delivering buffering ions of combinations or single component alkali metal polyoxylates (for example sodium silicate) to passivate the metal surface, e.g., refer to item 3) in the previous paragraph. In the case of sodium silicate, the carrier contains dissolved silica in the form of a silicate anion in water as well as sodium hydroxide in the presence of water. If desired, sodium hydroxide can be employed maintaining the pH of the solution in a range where the silicate can remain soluble. In the case of other substrates and other anion systems, the buffering capacity of the reactants is designed to passivate the surface, manage the pH of the surface chemistry, activate the surface, oxidize the surface, or to prepare or condition the surface for a mineral-forming reaction or any combination of the above. The delivery of ions is through a carrier comprising a membrane employing osmotic pressure to drive precursors to the surface.

20 The ionic species, which are present in the carrier or that pass through the carrier membrane, can then interact chemically and can become associated with the surface of the metal to form a submicron mineralization layer, e.g., a monolayer. In the present invention these

5 interactions occur adjacent to or upon the surface of the substrate to form a mineralized layer. It is to be understood that the aforementioned membrane is associated with creating an oxygen-limited passivation environment as part of the mineralization process.

Moreover, the mineralized layer precursors can interact in such a manner to produce mineralized layer in-situ at the surface. Depending upon the conditions of the surface, the substrate may contribute precursors in the form of metal ions. The metal ions of the substrate surface may exist as oxides, or the ions may have reacted with chemical species in the surrounding environment to form other metal species. In the case of a zinc substrate or surface, zinc can oxidize in the environment existing at the surface as zinc oxide, but may also form zinc carbonate from the exposure to carbon dioxide in the air. Under certain conditions, the zinc carbonate will predominate the surface species of the precursor to form the mineralized surface. In the case of other metal substrates, the ability of the surface to contribute ions to function as mineral precursors can be achieved by conditioning the surface, e.g., to populate the surface with oxide species that will participate as mineralized layer precursors.

10 To enhance mineral layer formation on at least a portion of the surface of a metal substrate, the metal surface may be prepared or pretreated. Metal surfaces normally tend to be covered with a heterogeneous layer of oxides and other impurities. This covering can hinder the effectiveness of the buffering and/or mineral layer formation. Thus, it becomes useful to convert the substrate surface to a homogenous state thereby permitting more complete and uniform mineral layer formation. Surface preparation can be accomplished using an acid bath to dissolve the oxide layers as well as wash away certain impurities. The use of organic solvents and detergents or surfactants can also aid in this surface preparation process. Phosphoric acid based cleaners, such as Metal Prep 79 (Parker Anchem), fall into a category as an example commonly used in industry. Other combinations of acids and cleaners are useful as well and are selected depending upon the metal surface and composition of the desired mineral layer. Once the surface is pretreated, the surface can then be subjected to further activation, if necessary, to enhance the buffering capability, including but not limited to oxidation by any suitable method. Examples of suitable methods comprise immersion in hydrogen peroxide, sodium peroxide, potassium permanganate, mixtures thereof, among other oxidizers.

5 Depending upon the carrier and process condition, precursors can pass through the carrier membrane system as anions, and interact adjacent to or upon the surface with metal cations, which in most cases are donated by the metal surface or substrate to form a relatively thin mineralized layer, e.g., a monolayer. In one aspect of the invention, sodium silicate (as SiO_3^- ion) reacts with a zinc containing surface, e.g., that exists primarily as zinc carbonate, to form an amorphous mineralization layer containing a nanocrystalline hemimorphite phase that is normally less than 100 Angstroms in thickness. In this aspect of the invention, the metal surface was prepared for mineralization by the presence of a suitable buffering alkali, e.g., buffering with a silicate to a pH in the range of about 9.5 to about 10.5. While a higher pH can be effectively used, a pH of less than about 11 minimizes the need for certain relatively complex and expensive handling procedures.

In another aspect of the invention, the delivery of pH buffering agents as well as the anion reactants can be designed to tailor the surface characteristic. For example, in order to achieve improved resistance to acid rain on a zinc surface, the silicate anions can be complemented with zirconate anions. Further, in the case of an iron containing surface, the carrier can deliver silicate anions to anodically conditioned surface to form amorphous phase comprising jujgoldite.

In yet another aspect of the invention, a surface pretreatment was used to enhance the mineralization layer formation. For example, in order to achieve an improved corrosion surface on iron, the steel substrate was first treated with a phosphoric acid containing cleaner, then exposed to an oxidizer in order to remove unwanted material and convert at least a portion of the pretreated metal surface to a homogeneous species of iron oxide. The mineralized layer was then formed on the pretreated metal surface by being contacted with sodium silicate containing precursor which in turn can proceed to form a clinopyroxene of sodium iron silicate.

While any suitable buffer can be employed for practicing the invention, buffer solutions are typically prepared by mixing a weak acid and its salt or a weak base and its salt. Acidic buffers, for example, can be prepared using potassium chloride or potassium hydrogen phthalate with hydrochloric acid of appropriate concentrations. Neutral buffers can be prepared by mixing potassium dihydrogen phosphate and sodium hydroxide, for example. Alkaline (basic) buffers can be prepared by mixing borax or disodium hydrogen phosphate with sodium hydroxide, for

5 example. Many more chemical combinations are possible, using appropriate chemicals to establish the proper sequence of proton transfer steps coupled with the intended reactions. Buffer exchange rates may be modified by combinations of buffer materials that react at different ionic exchange rates; buffers of low-change type react more rapidly than high-change types.

Aqueous polymers are preferred carriers for buffers in liquid form and include water-reducible aldehyds and modified aldehyds, acrylic latexes, acrylic epoxy hybrids, water-reducible epoxies, polyurethane dispersions, vinyl and ethylene vinyl acetates, and mixtures thereof. Such polymers are water vapor permeable but are repellent of liquid water and are essentially water insoluble after curing. These polymers can form a semipermeable membrane for water vapor and ionic transfer. Hence, if the surface of the metal substrate is dry, water vapor can permeate the membrane; but, buffering ions, which are present in the membrane or that pass through the membrane, can passivate the metal surface thereby reducing corrosion.

Buffer materials are chosen based on the type of the surface or substrate to be protected. Metal substrates may be protected from corrosion by passivating the substrate surface. Such passivation may generally be accomplished only in certain pH ranges which, in turn, depend on the 20 specific substrate to be protected. For example, iron based alloys are passivated with an alkaline pH (pH 8-12). This pH range is preferably accomplished with sodium silicate and/or potassium silicate powders; but other alkaline materials may be used. A blend of sodium and potassium silicates is also useful for achieving viscosity control in aqueous carrier/membrane formulations.

In a further aspect of the invention, a mineralized layer is obtained by mixing silicates and anodic oxidizing materials such as sodium carbonate and delivering the mixture in a manner effective to activate the metal surface.

While the above description emphasizes a zinc containing surface, the surface of a wide range of metal surfaces can be altered to impart beneficial surface characteristics. In most cases, the substrate or the surface thereof contributes cations to the mineralization-forming reaction. Examples of metal surfaces include aluminum, zinc, iron, copper, brass, iron, steel, stainless steel, lead, alloys thereof, among others. In the case of limited mineral layer formation or 30 thickness of the mineral layer is caused by the surface contribution of the cation, an improved result can be obtained by managing or tailoring the pH. That is, the buffering capacity and the pH of the carrier is substrate surface-specific and is tailored to manage the surface chemistry to

5 form the inventive mineralization layer, e.g., selecting a pH at which the surface is reactive and encourages formation of the mineralization layer. The reaction for forming the new surface with continue until such time that the finite quantity of metal atoms at the surface are consumed. If the new mineralized layer is melted or destroyed, a desirable aspect of the instant invention is that the surface will reinitiate mineralization formation with any available precursors. The ability to reinitiate mineralization or self-repair damaged surfaces is a novel and particularly desirable characteristic of the invention.

The delivery/method of the alkali metal polyoxolates (or mineralization layer) can be provided through a membrane from a reservoir as described in the U.S. Patent Application Serial No. 08/634,215; previously incorporated by reference. In the present invention, soluble precursors, such as silicate materials, are used within one or more coating layers. For example, in a polymer containing carrier system, one of the layers would be charged with sodium or potassium silicates wherein the outer layer(s) are employed to control the rate of moisture flow through the carrier. These carriers are typically relatively hard films as the normal polymerization of the carrier occurs to form a plastic type polymer type coating. Additional delivery methods have been developed utilizing soft films, gels, sealants, adhesives, and paints wherein the membrane feature is formed in-situ by the reaction between a silicate, e.g., sodium silicate, and silica. By controlling the quantity of the silica in the carrier, the mineralized layer can be designed to suit the specific application. Depending upon the pH and relative concentration of silicate and silica, the degree of crystal formation, e.g., a silica containing hemimorphite within an amorphous layer, can also be designed to achieve a predetermined result.

Without wishing to be bound by any theory or explanation the formation of the mineralized layer can occur under a wide range of conditions (normally ambient) and via a plurality of mechanisms. Normally, the mineralization layer forms underneath the carrier upon the metal surface, e.g., as buried layer under a carrier comprising a reservoir of precursors. If so formed, whatever ions are needed in the reservoir layer to form the mineralized layer, are expediently included as water soluble salts in the reservoir layer. On the other hand, all the ions employed to form the mineralized layer need not necessarily be included in the reservoir layer. That is, if desired cations can be supplied from the underlying metal surface and need not necessarily be

5 included as water soluble salts in the reservoir layer. Such cations can be obtained from the surface of the substrate metal itself, by reaction of the substrate with the anions of the precursor component for the mineralized layer. Since the mineralized layer is normally relatively thin, sufficient cations for the mineralized layer can even be supplied from the substrate when present only as an alloying ingredient, or perhaps even as an impurity. Additionally, the cations needed for the mineralized layer can be supplied from water soluble salts in the reservoir layer, as indicated above. Further, if the mineralized layer is to be formed from an overlying reservoir layer that also contains buffer components, at least some of the salts used for buffering can be employed for forming the mineralized layer. The latter reservoir layer would possess a self healing effect by functioning as a source of mineralization precursors in the event the layer was damaged. Once the mineralization layer has been formed to the degree desired, the carrier or reservoir layer can remain as a component of the finished article or removed.

In one embodiment of the invention, silicon can provide the predominate lattice-forming unit, e.g., CRN. The fundamental unit on which the structure of all silicates is based on consists of four O²⁻ at the apices of a regular tetrahedron surrounding and coordinated by one Si⁴⁺ at the center. A bond between silicon and oxygen atoms may be estimated by use of Pauling's electronegativity concept as 50% ionic and 50% covalent. That is, although the bond typically arises in part from the attraction of oppositely charged ions, the silicon-oxygen bond also involves sharing of electrons and interpretation of the electronic clouds of the ions involved. Each O²⁻ has the potential of bonding to another silicon atom and entering into another tetrahedral grouping, thus uniting the 10 tetrahedral groups through the shared (or bridging) oxygen. Such linking of tetrahedra is often referred to as polymerization. In the case of silicate polymerization, the amount of oxygen is normally less than the stoichiometric amount of 4:1, Si:O. Silicates in which two SiO₄ groups are linked can produce Si₂O₇ groups which are classed as disilicates. If more than two tetrahedra are linked, a closed ring like structure can be formed of a general composition Si_nO_{4n}. Fourfold tetrahedral rings have composition Si₄O₁₁. Tetrahedra may also be joined to form infinite single chains, called pyroxenes with a unit composition of SiO₃. Infinite double chains, classified as amphiboles, normally possess a ratio of Si:O = 4:11. Infinitely extending flat sheets are formed of unit composition Si₃O₈. Three dimensional framework silicates, such as feldspars or zeolites, typically result in a network oxide of unit composition SiO₂.

5 In the present invention, silicates can be polymerized into a wide range of Si:O ratios. In the case of a zinc containing surface, the surface can interact with a mineral layer precursor to form a silicate. In the case of an iron containing surface, the surface can interact with a suitable mineral layer precursor to form a chain silicate, e.g., a clinoperoxite. When silicate polymerization occurs in the instant invention, the covalency of the silicon-oxygen bond is decreased. This is evidenced by an increased binding energy of the photoelectron as detected by x-ray photoelectron spectroscopy, e.g., a binding energy between 102.0 to 103.3 eV in comparison to 101.5 eV for orthosilicate.

Without wishing to be bound by any theory of explanation it is also believed that the aforementioned inventive mineralized can comprise a complex oxide of the form:

15 M_xNyO_z , wherein "M" represents one or more cationic elements having a covalency factor of less than about 0.5 that functions to balance the charge of the complex oxide, "N" represents one or more lattice forming elements having a covalency factor greater than about 0.15 that functions as the structural component of the complex oxide and optionally wherein the NyO_z carries single or multiple crystal structures; and wherein x, y and z comprise any number the 20 total of which balances the charge of the complex oxide. In some cases, the covalency factor of M is less than about 0.33 and the covalency factor N is greater than about 0.33.

Applications for the films and coatings of the invention include, for example, components such as coatings and paints of components, parts and panels for use in the automotive industry, home-consumer products, construction and infrastructures, aerospace industry, and other out-door or corrosive applications wherein it is desirable to improve the characteristics of a metal surface and the use of heavy metals in elemental or non-elemental form is environmentally undesirable. The films and coatings may be applied to new products or over conventional platings to extend the useful service life of the plated component.

25 While the above description places particular emphasis upon forming a silicate containing mineralized layer, one or more mineralized layers having chemically similar or distinct compositions can be applied upon the same metal surface. If desired, the mineralized surface can be further modified by using conventional physical or chemical treatment methods.

The following Examples are provided to illustrate not limit the scope of the invention as defined in the appended claims. These Examples contain results of the following analysis: Auger

5 Electron Spectroscopy (AES), Electronic Impedance Spectroscopy (EIS), and Electron Spectroscopy for Chemical Analysis (ESCA) and Potentiodynamic (DC) Polarization. These analysis were performed using conventional testing procedures. The results of the AES demonstrate that the thickness of the mineralized layer can range from about 10 to 50 microns. EIS demonstrates that the mineralized layer imparts corrosion resistant properties to the surface, e.g., a reduced Icorr corresponds to a reduced corrosion current and in turn a reduced corrosion rate. The XPS data demonstrates the presence of a unique hemimorphite crystal within the mineralized layer, e.g., XPS measures the bond energy between silicon and oxygen atoms and compares the measured energy to standardized values in order to determine whether or not known crystals are present. Conventional X-ray diffraction analysis confirmed that the mineralized layer is predominately amorphous, e.g., an X-ray measurement resulted in wide bands thereby indicating the presence of an amorphous phase.

EXAMPLES

-Auger Electron Spectroscopy-

Examples 1-4 were prepared and analyzed in accordance with the following AES

20 procedure.

Instrumentation Used

Instrument:	Physical Electronics 545, Single Pass Cylindrical Analyzer
25 Sample Excitation:	Electron Gun
Electron Beam Detection:	Multiplier Voltage, 1 kV
Modulation Voltage, 3 eV	
30 Signal Detection:	Lock-in Magnification, 10X
Sputter Gun Settings: Beam Voltage, 3kV	Lock-in Time Constant, 0.001 sec.
Focus Dial Setting, 2	

5 Raster X Dial Setting, 10
 Raster Y Dial Setting, 10
 Emission Current, 15, mA
 Sputter Rate, 12 Å/min
 Start Time, 0 min.
 Final Time, 30 min
 Sputter Depth, 360 Å

Example 1

The coating had a first layer with the following formulation (by weight):

15 25% Water (Fisher Scientific)
 75% NeoRezR-9637 (Zeneca Resins)

and a second layer with the following formulation (by weight):

20 6.5% N-grade Sodium Silicate
 13% Water (Fisher Scientific)
 80.5% NeoRezR-9637 (Zeneca Resins)

25 The components were mixed by hand for approximately 15 minutes. The first layer was then

cast onto a standard 1010 steel test panel, obtained through ACT Laboratories, with a dry film thickness of about 0.5 to 0.7 mil. This layer was dried to tack free at 60 C for 10 minutes.
 30 The second layer was then applied, with a dry film thickness of about 0.5 to 0.7 mil. This layer was also dried to tack free at 60 C for 10 minutes. The coating was left on the panel for a minimum of 24 hours. Most of the coating was then removed with BIX Stripper (Walmart Stores) and a plastic spatula. The residual coating was washed off with copious amounts of Naphtha (Commercial Grade, Walmart Stores), and Reagent Alcohol (Fisher Scientific).

21

Example 2

The coating had one layer with the following formulation (by weight):

25 25% Water (Fisher Scientific)
 75% NeoRezR-9637 (Zeneca Resins)

30 and a second layer with the following formulation (by weight):

6.5% N-grade Sodium Silicate
 13% Water (Fisher Scientific)
 80.5% NeoRezR-9637 (Zeneca Resins)

The components were mixed by hand for approximately 15 minutes. The first layer was then cast onto a standard electrogalvanized test panel, obtained through ACT Laboratories, with a dry film thickness of about 0.5 to 0.7 mil. This layer was dried to tack free at 60 C for 10 minutes.
 20 The second layer was then applied, with a dry film thickness of about 0.5 to 0.7 mil. This layer was also dried to tack free at 60 C for 10 minutes. The coating was left on the panel for a minimum of 24 hours. Most of the coating was then removed with BIX Stripper (Walmart Stores) and a plastic spatula. The residual coating was washed off with copious amounts of Naphtha (Commercial Grade, Walmart Stores), and Reagent Alcohol (Fisher Scientific).

25 The coating had one layer with the following formulation (by weight):

25% Water (Fisher Scientific)
 75% NeoRezR-9637 (Zeneca Resins)

30 and a second layer with the following formulation (by weight):

22

5 6.5% N-grade Sodium Silicate
13% Water (Fisher Scientific)
80.5% NeoRezR-9637 (Zeneca Resins)

10 The components were mixed by hand for approximately 15 minutes. The first layer was then cast onto a standard zinc phosphated, 1010 steel test panel, obtained through ACT Laboratories, with a dry film thickness of about 0.5 to 0.7 mil. This layer was dried to tack free at 60 C for 10 minutes. The second layer was then applied, with a dry film thickness of about 0.5 to 0.7 mil. This layer was also dried to tack free at 60 C for 10 minutes. The coating was left on the panel for a minimum of 24 hours. Most of the coating was then removed with BIX Stripper (Walmart Stores) and a plastic spatula. The residual was washed off with copious amounts of Naptha (Commercial Grade, Walmart Stores), and Reagent Alcohol (Fisher Scientific).

Example 4

The coating had one layer with the following formulation (by weight):

20 25% Water (Fisher Scientific)
75% NeoRezR-9637 (Zeneca Resins)

and a second layer with the following formulation (by weight)

25 6.5% N-grade Sodium Silicate
13% Water (Fisher Scientific)
80.5% NeoRezR-9637 (Zeneca Resins)

30 The components were mixed by hand for approximately 15 minutes. The first layer was then cast onto a standard iron phosphated, 1010 steel test panel, obtained through ACT Laboratories, with a dry film thickness of about 0.5 to 0.7 mil. This layer was dried to tack free at 60 C for 10 minutes. The second layer was then applied, with a dry film thickness of about 0.5 to 0.7 mil. This layer was also dried to tack free at 60 C for 10 minutes. The coating was left on the panel

5 for a minimum of 24 hours. Most of the coating was then removed with BIX Stripper (Walmart Stores) and a plastic spatula. The residual was washed off with copious amounts of Naptha (Commercial Grade, Walmart Stores), and Reagent Alcohol (Fisher Scientific).

10 Each of the previously described washed test panels were passed through an AES analysis in accordance with conventional methods. This analysis confirmed the presence of the inventive mineral layer and generated data which demonstrated that for each substrate, the thickness of the mineralized layer was on the order of about 50 to about 70 Å thick.

Electrical Impedance Spectroscopy-

15 Examples 5 through 13 were prepared for Electrochemical Impedance Spectroscopy (EIS) analysis. EIS is one method of determining corrosion rates of a metal or a coated metal. In this technique, a small-amplitude sinusoidal potential perturbation was applied to the working electrode at a number of discrete frequencies ranging from 60,000 Hz to 0.0005 Hz. At each one of these frequencies, the resulting current waveform exhibited a sinusoidal response that was out of phase with the applied potential signal by a certain amount. The electrochemical impedance was a frequency-dependent proportionality factor that acts as a transfer function by establishing a relationship between the excitation voltage signal and the current response of the system. This method was detailed by the American Society for Testing and Materials (ASTM) in Electrochemical Corrosion Testing, STP 727.

Example 5
Example 5

A gel was prepared having the following formulation (by weight):

10% Cab-O-Sil TS-720 Silica
90% Amoco DURASYN™ 17/4 polyalphaolefin

15 The above formulation was mixed in a Hobart Mixer (model N-50) for approximately 30 minutes. The gel was then cast onto a standard electrogalvanized test panel, obtained through ACT Laboratories, at a thickness of 1/16" to 1/8". The gel was left on the panel for a minimum of 24 hours. Most of the gel was then removed with a plastic spatula. The residual gel was

5 washed off with copius amounts of Naptha (Commercial Grade, Walmart Stores), and Reagent Alcohol (Fisher Scientific).

Example 6

A gel was prepared having the following formulation (by weight):

10	10% Cab-O-Sil TS-720 Silica	10% Cab-O-Sil TS-720 Silica
	20% G-Grade Sodium Silicate (PQ Corporation)	20% Sodium Phosphate (Fisher Scientific)
	70% Amoco DURAST TM 174 polyalphaolefin	70% Amoco DURAST TM 174 polyalphaolefin

15 The above formulation was mixed in a Hobart Mixer (model N-50) for approximately 30 minutes. The gel was then cast onto a standard electrogalvanized test panel, obtained through ACT Laboratories, at a thickness of 1/16" to 1/8". The gel was left on the panel for a minimum of 24 hours. Most of the gel was then removed with a plastic spatula. The residual was washed off with copius amounts of Naptha (Commercial Grade, Walmart Stores), and Reagent Alcohol (Fisher Scientific).

20 This Example was repeated with the exception that sodium carbonate was employed instead of sodium phosphate.

Example 8
A gel was prepared having the following formulation (by weight):

10	10% Cab-O-Sil TS-720 Silica	10% Cab-O-Sil TS-720 Silica
	20% Sodium Phosphate (Fisher Scientific)	20% Sodium Phosphate (Fisher Scientific)
	70% Amoco DURAST TM 174 polyalphaolefin	70% Amoco DURAST TM 174 polyalphaolefin

15 The above formulation was mixed in a Hobart Mixer (model N-50) for approximately 30 minutes. The gel was then cast onto a standard electrogalvanized test panel, obtained through ACT Laboratories, at a thickness of 1/16" to 1/8". The gel was left on the panel for a minimum of 24 hours. Most of the gel was then removed with a plastic spatula. The residual was washed off with copius amounts of Naptha (Commercial Grade, Walmart Stores), and Reagent Alcohol (Fisher Scientific).

20 This Example was repeated with the exception that sodium carbonate was employed instead of sodium phosphate.

Example 7

A gel was prepared having the following formulation (by weight):

25	10% Cab-O-Sil TS-720 Silica	10% Cab-O-Sil TS-720 Silica
	20% Sodium Molybdate (Fisher Scientific)	20% Water (Fisher Scientific)
	70% Amoco DURAST TM 174 polyalphaolefin	75% NeoRezR-9637 (Zeneca Resins)

30 The above formulation was mixed by hand for approximately 15 minutes. The coating was then cast onto a standard 1010 steel test panel, obtained through ACT Laboratories, at a thickness for a total dry film thickness of 2.1 to 2.5 mils in three layers. Each layer was dried to tack free at 60 C for 1.5 minutes. The coating was left on the panel for a minimum of 24 hours. Most of the coating was then removed with BIX Stripper (Walmart Stores) and a plastic spatula. The residual was washed off with copious amounts of Naptha (Commercial Grade, Walmart Stores), and Reagent Alcohol (Fisher Scientific).

35 The above formulation was mixed in a Hobart Mixer (model N-50) for approximately 30 minutes. The gel was then cast onto a standard electrogalvanized test panel, obtained through ACT Laboratories, at a thickness of 1/16" to 1/8". The gel was left on the panel for a minimum of 24 hours. Most of the gel was then removed with a plastic spatula. The residual was washed off with copious amounts of Naptha (Commercial Grade, Walmart Stores), and Reagent Alcohol (Fisher Scientific).

5

Example 10

A coating was prepared having the following formulation (by weight):

6.5% N-grade Sodium Silicate	6.5% Sodium Molybdate Solution (274.21 grams/liter)
10 13% Water (Fisher Scientific)	10 13% Water (Fisher Scientific)
80.5% NeoRezR-9637 (Zeneca Resins)	80.5% NeoRezR-9637 (Zeneca Resins)

The above formulation was mixed by hand for approximately 15 minutes. The coating was then cast onto a standard 1010 steel test panel, obtained through ACT Laboratories, at a thickness for a total dry film thickness of 2.1 to 2.5 mils in three layers. Each layer was dried to tack free at 60 C for 15 minutes. The coating was left on the panel for a minimum of 24 hours. Most of the coating was then removed with BIX Stripper (Walmart Stores) and a plastic spatula. The residual was washed off with copious amounts of Naphtha (Commercial Grade, Walmart Stores), and Reagent Alcohol (Fisher Scientific).

20

Example 11

A coating was prepared having the following formulation (by weight):

6.5% Sodium Vanadate Solution (156.3 grams/liter)	6.5% Sodium Carbonate Solution (120.12 grams/liter)
25 13% Water (Fisher Scientific)	25 13% Water (Fisher Scientific)
80.5% NeoRezR-9637 (Zeneca Resins)	80.5% NeoRezR-9637 (Zeneca Resins)

The above formulation was mixed by hand for approximately 15 minutes. The coating was then cast onto a standard 1010 steel test panel, obtained through ACT Laboratories, at a thickness for a total dry film thickness of 2.1 to 2.5 mils in three layers. Each layer was dried to tack free at 60 C for 15 minutes. The coating was left on the panel for a minimum of 24 hours. Most of the coating was then removed with BIX Stripper (Walmart Stores) and a plastic spatula. The residual was washed off with copious amounts of Naphtha (Commercial Grade, Walmart Stores), and Reagent Alcohol (Fisher Scientific).

27

5

Example 12

A coating was prepared having the following formulation (by weight):

6.5% N-grade Sodium Silicate	6.5% Sodium Molybdate Solution (274.21 grams/liter)
10 13% Water (Fisher Scientific)	10 13% Water (Fisher Scientific)
80.5% NeoRezR-9637 (Zeneca Resins)	80.5% NeoRezR-9637 (Zeneca Resins)

The above formulation was mixed by hand for approximately 15 minutes. The coating was then cast onto a standard 1010 steel test panel, obtained through ACT Laboratories, at a thickness for a total dry film thickness of 2.1 to 2.5 mils in three layers. Each coat was dried to tack free at 60 C for 15 minutes. The coating was left on the panel for a minimum of 24 hours. Most of the coating was then removed with BIX Stripper (Walmart Stores) and a plastic spatula. The residual was washed off with copious amounts of Naphtha (Commercial Grade, Walmart Stores), and Reagent Alcohol (Fisher Scientific).

20

Example 13

A coating was prepared having the following formulation (by weight):

6.5% Sodium Vanadate Solution (156.3 grams/liter)	6.5% Sodium Carbonate Solution (120.12 grams/liter)
25 13% Water (Fisher Scientific)	25 13% Water (Fisher Scientific)
80.5% NeoRezR-9637 (Zeneca Resins)	80.5% NeoRezR-9637 (Zeneca Resins)

The above formulation was mixed by hand for approximately 15 minutes. The coating was then cast onto a standard 1010 steel test panel, obtained through ACT Laboratories, at a thickness for a total dry film thickness of 2.1 to 2.5 mils in three layers. Each layer was dried to tack free at 60 C for 15 minutes. The coating was left on the panel for a minimum of 24 hours. Most of the coating was then removed with BIX Stripper (Walmart Stores) and a plastic spatula. The residual was washed off with copious amounts of Naphtha (Commercial Grade, Walmart Stores), and Reagent Alcohol (Fisher Scientific).

28

5 and Reagent Alcohol (Fisher Scientific).

The cleaned samples from Examples 5 to 13 were then tested by EIS in accordance with the following method.

10 Instrument: Solartron 1287 Electrochemical Interface

Solartron 1260 Impedance/Gain-Phase Analyzer

ZWare and CorrWare Software by Scribner

Settings:

60,000 to 0.005 Hz.

10 steps/decade

5 mV rms AC signal

Test Solution: 1Molar Ammonium Sulfate, pH = 3.0 (using 0.1 N Sulfuric Acid) for electrogalvanized substrates.

1Molar Ammonium Sulfate, pH = 2.0 (using 0.1 N Sulfuric Acid) for

1010 Steel substrates.

20 B value: corresponds to the average of the Tafel slope $B = (B_{\text{corr}} - B_{\text{no corr}}) / (B_{\text{corr}} + B_{\text{no corr}})$

The following Table sets forth the results of the EIS Procedure for the zinc substrate in Examples 5-8.

EIS TABLE FOR ZINC SUBSTRATES

surface	B	i_{corr} $\mu\text{A}/\text{cm}^2$	ave. B	ave. i_{corr} $\mu\text{A}/\text{cm}^2$	EXPL	Comments
Bare Zinc	0.062632	193.3	0.067255	205.5	standard	
	0.060229	204.4			std	
	0.063586	226.3			std	
	0.101038	85.3			std	

¹ Cell leaked at the base, setting up a localized galvanic cell.

¹ Cell leaked underneath gasket.

ZINC EIS TABLE CONTINUED

5

surface	B (mV)	i_{corr} $\mu A/cm^2$	ave. B (mV)	ave. i_{corr} $\mu A/cm^2$	EXAMPL NO	Comments
Na_2MoO_4	0.052667	99.3	0.061923	128.5	7	
	0.072482	191.1		7		
	0.059900	98.5		7		
	0.065942	128.6		7		
	0.058622	125.0		7		
	0.098991	153.9		7		
Na_2PO_4	0.047173	95.3	0.049479	153.9	8	
	0.045828	141.0		8		
	0.053326	232.8		8		
	0.056754	178.1		8		
	0.0585574	194.0		8		
	0.035220	84.2		8		
Na_2CO_3	0.041397	68.4	0.047339	98.7	8	
	0.044320	121.8		8	7	
	0.046295	81.1		8		
	0.057343	123.4		8	3	

5

-	44	31.5	125	252
-	68	33.8	172	196, pH = 2.25
Steel - coating was not completely stripped	1	36.7	564	65
Steel	26	33.2	371	90
-	57	33.2	554	60
Steel - coating was restripped	1	42.4	32.6	1300
Steel	1	42.2	59.7	707
-	24	40.8	58.7	695
-	48	40.8	70.2	581, pH = 2

Coating Containing Sodium Vandate - Example 11

surface	Hours Immersed	B (mV)	R_p (ohm-cm ²)	i_{corr} ($\mu A/cm^2$)
steel	25	305.8	55.5	645
-	49	33.9	75.8	448
-	74	33.9	82.6	410, pH = 2.8
-	78	42.2	25.7	1640, pH = 2
-	1.5	50.5	70	721
-	21	36.5	92	397
-	44	33.3	126	265
-	68	33.2	161	2006, pH = 2.25
-	1	40.8	146	280
-	24	40.6	64.6	632
-	48	40.6	75.6	537, pH = 2

The results of the EIS indicates that the greatest corrosion resistance (low i_{corr}) for a zinc 10 substrate is obtained by a sodium silicate or sodium carbonate mineral layers.

The following Tables list the corrosion data for the steel surfaces of Examples 9-13.

² Cell leaked at the base, discarded this data point.

5 Coating Without Silicates-Example 9

surface	Hours	B (mV)	R _p (ohm-cm ²)	i _{corr} (μA/cm ²)
steel	24	39.4	68	580
-	49	33.8	55	614
-	74	33.8	116	290, pH = 2.86
-	99	33.8	25	1352, pH = 2.0
-	21	34.4	54	637
-	44	36.3	95.8	379
-	68	45	55.6	808, pH = 2.36
-	1	46.1	59.3	777
-	24	41.7	43.8	952
-	48	43.3	55.5	780, pH = 2.4
-	1	42	135	311
-	24.5	40.8	51.8	788
-	48	40.5	53	764, pH = 2.5

5 Coating Containing Sodium Molybdate - Example 12

surface	Hours	B (mV)	R _p (ohm-cm ²)	i _{corr} (μA/cm ²)
steel	21	37.4	51.3	729
-	44	44.4	45.6	973
-	68	41.4	54.3	763, pH = 2.35
steel - coating was not completely removed	1	37.7	190	198
steel	26	36.6	138	265
-	57	36.6	212	173
steel - coating was not completely removed)	1	33.3	112	353

Coating Containing Sodium Carbonate - Example 13

surface	Hours	B (mV)	R _p (ohm-cm ²)	i _{corr} (μA/cm ²)
steel	24	40.4	96.6	409
-	48	39.5	130	293, pH = 2.2
-	1	46.9	42.5	1109
-	24	48.2	28.6	1687
-	44	46	45.2	1018, pH = 2.15

surface	Hours Immersed	B (mV)	R _p (ohm·cm ²)	i _{corr} (μ A/cm ²)
steel	25	37.2	82.5	451
-	49	27.6	123.5	223
-	74	27.6	128.8	214, pH = 2.3
-	78	37.2	34.4	1080, pH = 2
-	1	41.8	56.8	735
-	24.5	44.2	33.5	1320
-	48	43.4	40.7	1066, pH = 2.9
-	1	39.5	112	353
-	24	40.6	96.6	409
-	48	38.1	130	293, pH = 2.2
-	1	45.7	46.7	978
-	24	46.5	29.7	1566
-	44	44.4	41.4	1072 pH = 2.15

Based upon the measured corrosion currents, sodium carbonate and sodium silicate are the most effective in reducing the iron thereby indicating a reduced corrosion rate.

- X-Ray Photoelectron Spectroscopy (XPS) -

X-ray Photoelectron Spectroscopy (XPS) was performed on a series of samples in accordance with conventional procedures in this art.

Instrumentation Used

Instrument: Physical Electronics 5701 LSci
 X-ray Source: Monochromatic aluminum
 Source Power: 350 watts
 Analysis Region: 2 mm X 0.8 mm
 Exit angle: 65°

35

5	Acceptance angle: ±7°
	Charge reference: B.E. of C-H,C) = 284.6 eV
	Charge neutralization: flood gun
	Sampling Depth: (3 λ) was 70 Å
10	

Example 14
 A coating was prepared having the following formulation (by weight):

25% Water (Fisher Scientific)
15% NeoRezR-9637 (Zeneca Resins)
15% NeoRezR-9637 (Zeneca Resins)
25% Water (Fisher Scientific)
75% NeoRezR-9637 (Zeneca Resins)

Example 15
 A coating was prepared having the following formulation (by weight):

25% Water (Fisher Scientific)
75% NeoRezR-9637 (Zeneca Resins)
25% Water (Fisher Scientific)
75% NeoRezR-9637 (Zeneca Resins)
75% NeoRezR-9637 (Zeneca Resins)

36

5 Most of the coating was then removed with BIX Stripper (Walmart Stores) and a plastic spatula. The residual was washed off with copius amounts of Naptha (Commercial Grade, Walmart Stores), and Reagent Alcohol (Fisher Scientific).

Example 16

10 A coating was prepared having the following formulation (by weight):

6.5% N-grade Sodium Silicate
13% Water (Fisher Scientific)
80.5% NeoRezR-9637 (Zeneca Resins)

15

The above formulation was mixed by hand for approximately 15 minutes. The coating was then cast onto a standard electrogalvanized test panel, obtained through ACT Laboratories, for a total dry film thickness of 2.1 to 2.5 mils in three layers. Each layer was dried to tack free at 60 C for 15 minutes. The coating was left on the panel for a minimum of 24 hours. Most of the coating was then removed with BIX Stripper (Walmart Stores) and a plastic spatula. The residual was washed off with copius amounts of Naptha (Commercial Grade, Walmart Stores), and Reagent Alcohol (Fisher Scientific).

Example 17

25 A coating was prepared having the following formulation (by weight):

6.5% N-grade Sodium Silicate
13% Water (Fisher Scientific)
80.5% NeoRezR-9637 (Zeneca Resins)

30

The above formulation was mixed by hand for approximately 15 minutes. The coating was then cast onto a standard electrogalvanized test panel, obtained through ACT Laboratories, for a total dry film thickness of 2.1 to 2.5 mils in three layers. Each layer was dried to tack free at 60 C for 15 minutes. The coating was left on the panel for a minimum of 24 hours. Most of the coating

37

38

5 was then removed with BIX Stripper (Walmart Stores) and a plastic spatula. The residual was washed off with copius amounts of Naptha (Commercial Grade, Walmart Stores), and Reagent Alcohol (Fisher Scientific).

Example 18

10 A coating was prepared having the following formulation (by weight):

6.5% N-grade Sodium Silicate
13% Water (Fisher Scientific)
80.5% NeoRezR-9637 (Zeneca Resins)

15

The above formulation was mixed by hand for approximately 15 minutes. The coating was then cast onto a standard electrogalvanized test panel, obtained through ACT Laboratories, for a total dry film thickness of 2.1 to 2.5 mils in three layers. Each layer was dried to tack free at 60 C for 15 minutes. The coated panel was then exposed to a post-cure heat treatment of 125° C for one hour, in a standard laboratory oven. The coating was left on the panel for a minimum of 24 hours. Most of the coating was then removed with BIX Stripper (Walmart Stores) and a plastic spatula. The residual was washed off with copius amounts of Naptha (Commercial Grade, Walmart Stores), and Reagent Alcohol (Fisher Scientific).

Example 19

10 A coating was prepared having the following formulation (by weight):

6.5% N-grade Sodium Silicate
13% Water (Fisher Scientific)
80.5% NeoRezR-9637 (Zeneca Resins)

The above formulation was mixed by hand for approximately 15 minutes. The coating was then cast onto a standard electrogalvanized test panel, obtained through ACT Laboratories, for a total dry film thickness of 2.1 to 2.5 mils in three layers. Each layer was dried to tack free at 60 C for 15 minutes. The coating was left on the panel for a minimum of 24 hours. Most of the coating

5 15 minutes. The coated panel was then exposed to a post-cure heat treatment of 175° C for one hour, in a standard laboratory oven. The coating was left on the panel for a minimum of 24 hours. Most of the coating was then removed with BIX Stripper (Walmart Stores) and a plastic spatula. The residual was washed off with copious amounts of Naphtha (Commercial Grade, Walmart Stores), and Reagent Alcohol (Fisher Scientific).

10

Example 20

For purposes of comparing the results achieved by Examples 14-19, an electrogalvanized panel was soaked for 24 hours in a solution that had the following formulation (by weight):

15 25 mg/l Sodium Silicate

The sample panel was allowed to air dry.

Example 21

20 For purposes of comparing the results achieved by Examples 14-19, an electrogalvanized panel was soaked for 24 hours in a solution that had the following formulation (by weight):

25 mg/l Sodium Silicate

25 The sample panel was allowed to air dry. The panel was then exposed to a post-dry heat treatment of 125° C for one hour, in a standard laboratory oven.

Example 22

30 For purposes of comparing the results achieved by Examples 14-19, an electrogalvanized panel was soaked for 24 hours in a solution that had the following formulation (by weight):

25 mg/l Sodium Silicate

5 The sample panel was allowed to air dry. The panel was then exposed to a post-dry heat treatment of 175° C for one hour, in a standard laboratory oven.

Example 23

A crystal sample of Zn_2SiO_4 , which was air fractured and immediately introduced into the sample chamber of the XPS. The total air exposure was less than 2 minutes. The surface was examined initially with a low resolution survey scan to determine which elements were present. High resolution XPS spectra were taken to determine the binding energy of the elements detected in the survey scan. The quantification of the elements was accomplished by using the atomic sensitivity factors for a Physical Electronics 5701 LSci ESCA spectrometer.

10 The following Table sets forth the silicon binding energies which were measured for Examples 14-23.

Example Number	Treatment	Si 2p peak Energy (eV)	
		Room Temp	101.9
14	Room Temp	101.8	101.8
15	125 °C		
16	Room Temp	102.8	
17	Room Temp	102.8	
18	125 °C	102.6	
19	175 °C	102.6	
20	25 ppm Na_2SiO_4 , 25 °C	101.7	101.7
21	25 ppm Na_2SiO_4 , 125 °C	101.7	101.7
22	25 ppm Na_2SiO_4 , 175 °C	101.8	101.8
23	Zn_2SiO_4 crystal	101.5	101.5

The above Table illustrates that there has been a change in the concentration of Si as well

5 as the bond energy thereby providing further confirmation of the presence of the mineralized layer. Further, the Si-O bond energy of conventional zinc silicate crystal as well as the silicate soaked panels is distinct from the inventive mineralized layer.

Example 24

10 A coating was prepared having the following formulation (by weight):

6.3% N-grade Sodium Silicate
13% Water (Fisher Scientific)
80.5% NeoRezR-9637 (Zaneca Resins)

15 The formulation above was mixed by hand for approximately 15 minutes. The coating was then cast onto a standard electrogalvanized test panel, obtained through ACT Laboratories, for a total dry film thickness of 2.1 to 2.5 mils in three coats. Each coat was dried to tack free at 60°C for 15 minutes. The panel was exposed to ASTM B117 Salt Fog Chamber for 2400 hours. At the end of the salt fog exposure, there were large areas of the panel that were uncorroded. The area of uncorroded surface was cut into a small square sample and the urethane coating was removed by hand, using small tweezers. The sample was analyzed in accordance with the previously identified XPS method on an instrument of comparable sensitivity and accuracy. The results of the XPS are set forth in the following Table.

25 Example 24 Silicate Containing Coating Followed by Salt Spray Exposure

Example Number	Treatment	Si 2p peak Energy (eV)	Relative Concentration of Silicon on the Surface (% wt)
14	urethane coating, room temp	101.9	negligible at 1.0 %
15	urethane coating, 125 °C	101.8	negligible at 0.8 %

25

16	silicate, room temp	102.8	19.8 %
17	silicate, 125 °C	102.8	18.9 %
18	silicate, room temp	102.6	15.8 %
19	silicate, 175 °C (heat applied after coating removed)	102.6	11.6 %
20	immersion, 25 ppm Na ₂ SiO ₄ , 25 °C	101.7	4.1 %
21	immersion, 25 ppm Na ₂ SiO ₄ , 125 °C	101.7	3.2 %
22	immersion, 25 ppm Na ₂ SiO ₄ , 175 °C	101.8	3.9 %
23	Zn ₂ SiO ₄ crystal	101.5	NA
24	silicate, room temp, 2400 hours of B117 exposure	102.6	8.1 %

5

The above Table illustrates that the binding energy for the surface exposed to the silicate containing coatings range from 102.6 to 102.8. By observation of the XPS peak, it was observed that these energy values were actually manifold values from manifold peaks that contain more than one material. These materials can be characterized as a disilicate, or hemimorphic (bonding energy at 102.2), altered by the presence of Si-O bonds, as in SiO₂ (bonding energy at 103.3) and Si-O-C bonds with a bonding energy of 103.6 or 103.7. Such a bonding energy and

5 concentration of Si on the surface are distinct from conventional silica or silicate structures. The binding energy, for zinc silicate, Zn_2SiO_4 , 101.5 eV, is also distinct from the binding energy of the mineralized layer, 102.7 eV, or the hemimorphite.

EXAMPLE 25

10 This Example illustrates a process for pretreating a metal surface in order to enhance formation of the inventive silicate containing layer/surface.

1. Immersc panel in solution of 25% Metalprep 79 (Parker-Amchem) for 2 minutes,
2. Remove Panel and rinse with deionized water,
- 15 3. Scrub panel gently with "Kim-Wipe",
4. Rinse with deionized water and wipe off excess with Kim-wipe,
5. Immersc panel in 0.1 M NaOH solution for 10 sec.,
6. Wipe off excess with Kim-wipe,
7. Immersc panel in 30% H₂O₂ solution for 5 min.; and,
- 20 8. Wipe off excess with Kim-wipe.

EXAMPLE 26

A cold rolled steel panel (ACT Labs) was prepared with the pretreatment process described in Example 25. The pretreated panel was then coated with a first layer comprising the following formulation (by wt%):

- 25 25% Water (Fisher Scientific); and,
75% NeoRezR-9637 (Zeneca Resins).

30 A second and third layer comprising the following formulation were then applied upon the first layer (by weight):

13% N-grade Sodium Silicate,

Each formula was mixed by hand for approximately 10 minutes. Each layer was applied at a 1.2 mil wet film thickness and given a 15 minute 60 C cure to allow for a tack-free finish. The

5 6.5% Water (Fisher Scientific); and,
80.5% NeoRezR-9637 (Zeneca Resins).

10 Each aforementioned formulation was mixed by hand for approximately 10 minutes.

15 Each layer was applied at a 1.2 mil wet film thickness and cured for 15 minutes at 60 C to form for a tack-free finish. The panels were allowed to set overnight (or longer) and the coating was physically removed by hand.

20 XPS analysis was performed on the surface of this panel. The characteristic NaFe₃(SiO₃)₂Si(Cp) photoelectron binding energy was used to identify the species on the surface. The surface was characterized as a iron pyroxene species with the empirical formula of NaFe₃(SiO₃)₂. Comparison of a clinopyroxene, a naturally occurring iron pyroxene, indicates that the species on the surface of the panel is indeed a matrix of a polymerized SiO₃ and iron.

EXAMPLE 27

25 A electrozinc galvanized panel (ACT Labs) was prepared with the pretreatment method described above in Example 25. The panel was then coated with the following first formula (by wt%):

25% Water (Fisher Scientific); and,
75% NeoRezR-9637 (Zeneca Resins).

30 A second and third layer with the following formulation (by weight) were applied upon the first layer:

6.5% N-grade Sodium Silicate,
13% Water (Fisher Scientific); and,
80.5% NeoRezR-9637 (Zeneca Resins).

5 panels were allowed to set for 1 hour and the coating was physically removed by hand.

EXAMPLE 28

A electrozinc galvanized panel (ACT Labs) was prepared with the pretreatment method described in Example 25. The panel was then coated with two layers of the following formula (by wt%):

1.0% N-grade Sodium Silicate,
18.5% Water (Fisher Scientific); and,
80.5% NeoRezR-9637 (Zeneca Resins).

15 Each formula was mixed by hand for approximately 10 minutes. Each layer was applied at a 1.2 mil wet film thickness and given a 15 minute 60 C cure to allow for a tack-free finish. The panels were allowed to set for 24 hours and the coating was physically removed by hand. The coated galvanized panels were recovered and analyzed in accordance with the previously described ESCA/XPS methods. XPS analysis was performed on both panels. A first panel shows the Si(2p) photoelectron binding energy of 102.1 eV representing a zinc disilicate species. A second panel also shows the same binding energy at 102.1 eV also indicating the presence of a zinc disilicate species on the surface of the zinc. The second test panel also has significantly more silica on the surface, represented by the 103.3 eV binding energy than does the first panel. Because of the accumulation of silica on the surface of panel #2, the relative amount of zinc decreases due to the limited sampling depth of XPS. The Zn/Si ratio goes down from 2.0 to 0.43 on panel #1 to panel #2, respectively, as would be expected when the build up of silica increases on the panel surface. In both cases, the formation of a zinc disilicate protective species was detected.

30

EXAMPLE 29

Cold rolled steel panels (ACT Laboratories) were coated with the following formula (by wt%) to form a first layer:

5 25% Water (Fisher Scientific); and,
75% NeoRezR-9637 (Zeneca Resins).

A second layer with the following formulation (by wt%) was applied upon the first layer:

10 X% N-grade Sodium Silicate,
(19.5-X)% Water (Fisher Scientific); and,
80.5% NeoRezR-9637 (Zeneca Resins),
15 wherein X is a number, either 1.0 or 0.01 wt%, as described below. A second layer with the same composition of the first layer was applied. Each formula was mixed by hand for approximately 10 minutes. Each layer was applied at a 1.2 mil wet film thickness and given a 15 minute 60 C cure to allow for a tack-free finish. Each panel was given a heat treatment for 1 hour at the temperature described in the table below. The panels were allowed to set overnight.

	Panel	Silicate conc Set # (X%)	Heat temp (C)	B2247 fail time	Adhesion rating
20	1	0.01	60	8	1
	2	0.01	125	8	9
	3	1.0	60	24	1
	4	1.0	125	24	10

Two sets of tests were performed to examine the degree of protection against corrosion and the adhesion properties of the urethane coating to the steel surface. Corrosion protection was measured by time to reach 5% red rust coverage by humidity exposure according to ASTM B2247. Adhesion was measured by immersing the panels in deionized water for 10 minutes followed by 24 hour humidity exposure. The panel coatings were then scribed using a razor blade in a "X" pattern. Tape was placed over the scribe and pulled back. Performance was rated on a scale of 1-10, 1 = no adhesion and complete removal of coating, 10 = excellent adhesion, no

5 removal of coating.

The results, illustrated in the table above, show that increased loadings of silicate in the urethane coating improve corrosion protection. For best results, the amount of mineral layer precursor or sodium silicate is greater than 0 and less than about 7 wt.%, normally about 0.1 to about .01 wt% silicate, and heat treated at a temperature of about 125 C.

It is to be understood that the foregoing is illustrative only and that other means and techniques may be employed without departing from the spirit or scope of the invention as defined in the following claims.

5

The coatings formulations were used to coat electrogalvanized steel panels and hot-dipped galvanized zinc panels. The electrogalvanized panels were supplied by ACT (ACT E60 EZG 60G) 2 side, clean, unpolished. The hot-dipped galvanized panels were obtained from a metal building supplier (Bulter) and cut to size specifications. Each coating layer was applied with a #12 Ir. Drawdown rod to produce a 1.2 mil wet film coating thickness. Each coating layer was dried to a tack free condition by baking in a forced air convection oven at 60 C for 15 minutes before subsequent coating layers were applied. The overall dry film coating thickness on all of the panels averaged 1.4 mils.

15

EXAMPLE 30

The corrosion resistance of the following formulations were evaluated by salt spray testing per ASTM-B117 specifications with the panels positioned with the 6 inch long edge at the top and the bottom to preclude each of the test areas on each panel from affecting the adjacent areas.

20

The coating formulations were prepared as follows:

1. Formula #1: was prepared by adding 781 g. water to 2342 g. NeoRez R9637 polyurethane dispersion (Zeneca Resins) to achieve a composition of 75% by weight R9637 and 25% by weight water. The viscosity of the composition was 35 cp Brookfield (#2 spindle, speed 20, 70°F).

25

2. Formula #2: was prepared by diluting 143.5 mL N grade sodium silicate solution (PQ Corporation) with 345 mL distilled water and slowly mixing this into 2134.5 grams of NeoRez R9637 resin while stirring with an air powered Jiffy Mixer for approximately 15 minutes. The viscosity of the composition was 38 cp Brookfield (#2 spindle, speed 20, 70°F) and had a pH of 10.5 (pH paper).

5 The coatings formulations were used to coat electrogalvanized steel panels and hot-dipped galvanized zinc panels. The electrogalvanized panels were supplied by ACT (ACT E60 EZG 60G) 2 side, clean, unpolished. The hot-dipped galvanized panels were obtained from a metal building supplier (Bulter) and cut to size specifications. Each coating layer was applied with a #12 Ir. Drawdown rod to produce a 1.2 mil wet film coating thickness. Each coating layer was dried to a tack free condition by baking in a forced air convection oven at 60 C for 15 minutes before subsequent coating layers were applied. The overall dry film coating thickness on all of the panels averaged 1.4 mils.

10

15 Sample 1:

Sample Name: Equal Control Sample

Substrate: ACT E60 EZG 60G 2 Side E-Galv., clean, unpolish zinc coated panel.

Coating: None (control sample)

Additional Treatments: None

20 Sample 2:

Sample Name: Hot-dipped Control Sample

Substrate: HotDipped Zinc Panel 03X06X075

Coating: None

Additional Treatments: None

25 Sample 3:

Sample Name: Urethane Control Sample

Substrate: ACT E60 EZG 60G 2 Side E-Galv., clean, unpolish zinc coated panel.

Coating: 3 coats of Formula #1

Additional Treatments: None

30 Sample 4:

5 Sample Name: Heated Urethane Control Sample
 Substrate: ACT E60 EZG 60G 2 Side E-Galv., clean, unpolish zinc coated panel.
 Coating: 3 coats of Formula #1
 Additional Treatments: 75°C for 1 hour

10 Sample 5:
 Sample Name: Single Formula #2 Coat
 Substrate: ACT E60 EZG 60G 2 Side E-Galv., clean, unpolish zinc coated panel.
 Coating: 1 coat of Formula #2
 Additional Treatments: None

15 Sample 6:
 Sample Name: Single Formula #2 + Topcoat
 Substrate: ACT E60 EZG 60G 2 Side E-Galv., clean, unpolish zinc coated panel.
 Coating: 1 coat of Formula #2, than 1 coat of Formula #1
 Additional Treatments: None

20 Sample 7:
 Sample Name: 2 layers of Formula #2
 Substrate: ACT E60 EZG 60G 2 Side E-Galv., clean, unpolish zinc coated panel.
 Coating: 2 coats of Formula #2
 Additional Treatments: None

25 Sample 8:
 Sample Name: Single layer of Formula #2 + Topcoat
 Substrate: HotDipped Zinc Panel
 Coating: 1 coat of Formula #2, than 1 coat of Formula #1
 Additional Treatments: None

5 Sample 9:
 Sample Name: 3-layer Formula #2
 Substrate: ACT E60 EZG 60G 2 Side E-Galv., clean, unpolish zinc coated panel.
 Coating: 3 coats of Formula #2
 Additional Treatments: None

10 Sample 10:
 Sample Name: 3-layers of Formula #2
 Substrate: HotDipped Zinc Panel
 Coating: 3 coats of Formula #2
 Additional Treatments: None

15 Sample 11:
 Sample Name: 2 layers of Formula #2 + Topcoat
 Substrate: ACT E60 EZG 60G 2 Side E-Galv., clean, unpolish zinc coated panel.
 Coating: 2 coat of Formula #2, than 1 coat of Formula #1
 Additional Treatments: None

20 Sample 12:
 Sample Name: 2 layers of Formula #2 + Topcoat
 Substrate: HotDipped Zinc Panel
 Coating: 2 coat of Formula #2, than 1 coat of Formula #1
 Additional Treatments: None

25 Sample 13:
 Sample Name: 2 layers of Formula #2 + Topcoat + Heat
 Substrate: ACT E60 EZG 60G 2 Side E-Galv., clean, unpolish zinc coated panel.
 Coating: 2 coat of Formula #2, than 1 coat of Formula #1
 Additional Treatments: 75° C for 1 hour

5

Sample 14:

Sample Name: 2 layers of Formula #2 + Topcoat + Heat

Substrate: ACT E60 EZG 60G 2 Side E-Galv., clean, unpolish zinc coated panel.

Coating: 2 coat of Formula #2, then 1 coat of Formula #1

Additional Treatments: 175° C for 1 hour

The results of analyzing the above Samples in accordance with ASTM B117.

No.	Sample	Substrate	Performance Hours ASTM B117		Rust
			1st Red	5% Red	
1	Zinc E-Galv Control		24	120	
2	Zinc Hot-dipped Control		24	528	
3	Urethane Control		320	452	
4	Heated Urethane Control		344	672	
5	Single Layer		864	1080	
6	Single Layer + Topcoat		876	1100	
7	2-Layer		1288	1476	
8	Single Layer + Topcoat		1308	1500	
9	3-Layer		1206	1320	
10	3-Layer		1200	1300	
11	2-Layer + Topcoat		1336	1608	
12	2-Layer + Topcoat		2000	2400	
13	2-Layer + Topcoat + Heat		2300+	2600+	
14	2-Layer + Topcoat + Heat		2500+	3000+	
15					30

15 The above Table illustrates that the instant invention can be employed for increasing the corrosion resistance of a zinc containing surface by a factor of at least 8, e.g., compare samples 4

51

Example 31
This Example illustrates whether or not corrosion resistance is imparted by an alkaline pH (attributed to Sodium Hydroxide) and whether the presence of the soluble silicate ion offers any additional protection beyond the contribution of the alkaline pH.

This Example was performed by preparing and testing three sets of formulations:

1. 3 layers of polymer with no additives.
2. 3 layers of water-borne polymer with a pH adjusted to that of a alkali silicate doped coating.
3. 3 layers of polymer with sodium silicate added.

The coating formulations were prepared as follows:

1. The coating for formulation 1. was prepared by adding 781 g. water to 2342 g. NeoRez R9637 polyurethane dispersion (Zeneca Resins) to achieve a composition of 75% by weight R9637 and 25% by weight water. The viscosity of the composition was 35 cP Brookfield (#2 spindle, speed 20, 70°F).

2. The coating for formulation 2. was prepared by dissolving 2.4335g. NaOH (sodium hydroxide powder) in 41.2 G. of distilled water. This sodium hydroxide solution was slowly mixed into 254 g. NeoRez R9637 resin with an air powered Jiffy Mixer over the course of 5 minutes. The viscosity of the composition was 60 cP Brookfield (#2 spindle, speed 20, 70°F) and had a pH of 10.5 (pH paper).

3. The coating for formulation 3. was prepared by diluting 143.5 mL N grade sodium silicate solution (PQ Corporation) with 345 mL distilled water and slowly mixing this into 2134.5 grams of NeoRez R9637 resin while stirring with an air powered Jiffy Mixer for approximately 15 minutes. The viscosity of the composition was 38 cP

5 Brookfield (#2 spindle, speed 20, 70°F) and had a pH of 10.5 (pH paper).

The coating formulations were used to coat electrogalvanized steel panels. The electrogalvanized panels were ACT E60 EZG 60G 2 side, clean, unpolished from APR 29396 Batch 30718614.

Each coating layer was applied with a #12 Jr. Drawdown rod to produce a 1.2 mil wet film convection oven at 60 C for 15 minutes before subsequent coating layers were applied. The overall dry film coating thickness on all of the panels averaged 1.4 mils. The following panels were prepared:

surface	First Layer	Second Layer	Third Layer	alkali
Egolv	Formula #1	Formula #1	Formula #1	PANTST27
Egolv	Formula #3	Formula #3	Formula #3	ConNaOH
Egolv	Formula #2	Formula #2	Formula #2	Zn-NaOH

15

Five panels each having the surface and layering shown in the above Table were subjected to ASTM-B117 salt spray exposure until all panels in the group developed red corrosion on 5% of its surface, or approximately 1500 hours of test exposure was achieved. The average time in test hours to the development of first red corrosion and to development of red corrosion on 5% of each sample surface(Failure) is shown in the Table below:

AVERAGE SALT SPRAY RESULTS FOR ELECTROGALVANIZED STEEL PANELS

Coating Formulation	Avg Hours First Red	Avg Hours 5% Red (Fail)
Polyurethane Only	308	448

25

*Two test panels did not develop red corrosion during the ASTM B117 Test.
**Two test panels did not develop red corrosion during the ASTM B117 Test.

Polyurethane pH Adjusted	456	616
Polyurethane + Na Silicate	1176*	1336**

5 *Two test panels did not develop red corrosion during the ASTM B117 Test.

**Two test panels did not develop red corrosion during the ASTM B117 Test.

The above two Tables illustrate that when applied over electrogalvanized steel panels, polyurethane coatings which contain sodium silicate perform a minimum of three times better than the polyurethane coating by itself, whereas only a 1.5 times improvement can be obtained by adjusting the polyurethane coating pH to 10.5. The presence of the sodium silicate imparts a corrosion resistance benefit significantly greater than that arising from the alkaline pH.

Example 32

The purpose of this Example is to determine the effect on corrosion resistance of adding different loadings of G grade sodium silicate powder (PQ Corporation) to a urethane carrier. The 1.5,10,15,20,25,30 wt. % sodium silicate batches were prepared by adding 2,10,20,30,40,50, and 60 grams, respectively, of G grade sodium silicate powder(PQ Corporation) to 200 grams of PAO Grease(AM940126 supplied by Nye Lubricants Inc.) and mixing by hand for approximately 30 minutes.

The mixed silicate containing grease was evaluated by applying the grease upon ACT EZG 60G 2 Side E-Galv, clean, unpolish zinc coated panels. The grease was applied to the panels by applying an excess to the panels and running a gate type applicator across the panels to leave behind a 1/16 inch thick gel coating. Two panels were coated per substrate/grease loading.

20 Prior to testing, the grease was removed from the bottom 3-3.5 inches of the panels by lightly scraping the excess off with manilla tags, leaving a thin film of gel on the surface. The base oil of the gel was then removed from this thin film by soaking and rinsing with naphtha. The cleaned surface of the panels appeared to be covered with a thin white deposit postulated to be silica and sodium silicate. Each test panel consisted of an uncoated area approximately 1 inch wide at the hole end of the panel followed by a 1.5-2 inch width containing unremoved gel and finally containing a 3-3.5 inch wide area where the gel had been removed.

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The panels were evaluated by salt spray testing per ASTM-B117 specifications with the panels positioned with the 6 inch long edge at the top and the bottom to preclude each of the test areas on each panel from affecting the adjacent area.

		Salt Spray Hours (ASTM B117)		
test basis	silica	Panel 1	Panel 2	Average hours
0	First Red	480	672	576
	5% Red	648	1224	936
1	First Red	504	816	660
	5% Red	792	1248	1020
5	First Red	1296	1320	1308
	5% Red	2064	1872	1968
10	First Red	1320	1368	1344
	5% Red	1584	1536	1560
15	First Red	1176	1176	1176
	5% Red	1416	1536	1476
20	First Red	768	1200	984

The corrosion resistance of zinc coated panels was affected by the amount of silicate in the carrier. The above Table demonstrates that the presence of at least 1% sodium silicate significantly enhances the corrosion performance.

EXAMPLE 33

Electrozing galvanized steel panels (ACT Laboratories) were coated with two layers of the following formula (by wt%)

wherein X is a number, either 1.0, 0.1, or 0.01, as described below in greater detail. A final layer with the same composition of the first layer was applied. Each formula was mixed by hand for approximately 10 minutes. Each layer was applied at a 1.2 mil wet film thickness and given a 15 minute 60°C cure to allow for a tack-free finish. The panels were allowed to set overnight.

Two sets of tests were performed on these panels to examine the degree of formation of a zinc disulfide protective species and the adhesion properties. XPS analysis was performed on the panels with the urethane layer removed. The Si(2p) photoelectron binding energy signature

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5 of 102.1 eV representing the zinc disilicate protective species was examined to determine its presence. Adhesion was measured by immersing the panels in deionized water for 10 minutes followed by 24 hour humidity exposure. The panel coatings were then scribed using a razor blade in a "X" pattern. Tape was placed over the scribe and pulled back. Performance was rated on a scale of 1-10, 1= no adhesion and complete removal of coating, 10= excellent adhesion, no removal of coating.

Panel	Silicate conc (X%)	Formation of Zinc Disilicate	Adhesion rating
1	1.0	Yes	10
2	0.1	Yes	10
3	0.01	No	10

10 All panels show excellent adhesion of the urethane layer to the surface of the zinc substrate. A silicate concentration of about 0.01 wt% in the urethane formula used above does not allow for the formation of zinc disilicate on the surface of the zinc.

15 This Example illustrates the partial mineral formation on zinc galvanized surfaces. The zinc surface was prepared under controlled conditions to capture the formation process at different stages and was analyzed using ESCA, AFM, and salt spray exposure in order to correlate formation to salt spray performance. Testing illustrated that there is a correlation between hemimorphic formation and salt spray performance. The data also shows a possible reaction pathway.

20 The purpose of this experiment was to control the formation of the zinc silicate species on the surface. When the protective species possess the characteristic peak, the intensity should correlate to the degree of corrosion protection. The intensity of the peak was affected by the thickness of the zinc silicate formed and/or the percent coverage of the surface. Thicker coverage prolonged corrosion protection however, incomplete coverage produced immediate failure. To slow down the mineral layer formation rate on the surface, three techniques were employed. The

5 first was to lay down a "non-active" urethane coat (one without any silicate). When the active layer is applied, the silicate will then have a barrier of diffusion to migrate through slowing down the rate of reaction. The next technique was to apply active coatings with lower amounts of silicate done under the assumption the reaction rate is dependent upon this reactant. The last method was to remove the coating at different times, thus stopping the reaction at known times.

10 Electrozinc galvanized panels were obtained from ACT Laboratories (EZG 60G). Each panel was washed with reagent alcohol to remove any oils. The panels were coated based on the scheme showed below in table 1. The three factors being studied are the number of base coats (0 or 1), the concentration of the silicate used in the urethane layer (1% or 6.5%), and time of coating removal (1 hr or 2 days). Each factor and the recipe matrix is shown in table 1. Each panel will be coated with or without the base coat as called for by the experimental matrix. The panel was then be coated with two layers of the active coat using the specified concentration.

15 Coating was done using a #12 draw down bar. Each coat was cured for 15 minutes at 60 C. The removal of the coating was done by applying an excessively thick coating of urethane. This was done to increase tensile strength of the coating to facilitate the easy removal of the coating.

20 The base resin used for all the coatings was NeoRez R9637 (Zeneca). Resin concentration was held at 80.5 wt%. The appropriate amount of N-grade sodium silicate solution (PQ) was diluted and slowly added into the resin while stirring. Concentrations of the silicate solutions before dilution were 0.1, or 6.5 wt%. The solution was then topped off with water.

Samples were tested in accordance with the previously described ESCA analysis techniques. The results of the ESCA testing are set forth in the following table.

Panel	Strategy	Si(2p)	Species	Zn:Si

4	0-20	103.2	silica	0.26
	48hrs, 6.5%			
3	0-20	102.0 (60%)	disilicate	(0.4± 0.1)
	48hrs, 1%	103.1 (40%)	silica	0.1
2	1-20	102.1 (83%)	disilicate	1.07
	48hrs, 6.5%	103.3 (17%)	silica	
1	1-20	101.6	orthosilicate	2.63
	2hrs, 6.5%			
Willemite		101.3	orthosilicate	2.0
Hemimorphite		101.8	disilicate	2.0

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The Table shows the progression of formation on the zinc surface. Panel 1 shows the initial formation of a zinc orthosilicate species. As time progresses, the zinc disilicate became the predominant species on the surface as seen in panel 2. Also the presence of silica was seen in relatively low yields. As the reaction progresses, the relative amount of silica increased until it overshadows the whole spectra as seen in panel 4. The increasing amount of silica does not indicate a lower yield of zinc disilicate. The ESCA analysis is limited to the top 50 angstroms of a surface. The trend of the zinc to silicon ratio illustrated that there was an accumulation of silica as the ESCA did detect the zinc due to the increased levels of silica. Natural examples of orthosilicate and disilicate known as Willemite and Hemimorphite respectively, were tested for purposes of comparison. The binding energies show a 0.3 eV shift compared to the test panels. This was due to the presence of sodium ions which has been detected by ESCA. The presence of sodium ions decreased the covalency of the Si-O bond thus increasing the binding energy.

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EXAMPLE 35
The purpose of this Example is to demonstrate using potentiodynamic (DC) polarization to measure corrosion resistance of a mineralized layer.

The initial corrosion resistance on treated metal substrates (as described below in greater detail) were measured via the following electrochemical polarization method. The tests were performed utilizing a Solartron SI 1287 electrochemical interface connected to an electrochemical cell similar to that shown in ASTM-G5-87. The working electrode consisted of a 2 cm x 7.5 cm test panel held by a stainless steel alligator clip with copper wire attached to the alligator clip. The assembly was maintained in a stationary position by holding the alligator clip tight against the end edge of a 13 mm inside diameter glass tube with the wire attachment running up the inside of the tube and the clamping lever arm extending outside of the tube. The tube was fitted to the glass flask by running it through a rubber stopper and the wire was held tight within the glass tube by running it through a rubber stopper at the top end of the glass tube. The sample to be tested was positioned so that the alligator clip connection to the test panel does not contact the cell test solution. The counter electrode was a standard platinum electrode and the reference electrode was a standard saturated calomel electrode connected to the test cell via a salt bridge. Tests were performed at ambient lab conditions of 70 F.

The DC polarization tests were performed by adding 700 mL of 5.0 weight % sodium chloride solution to the flask and aeration vigorously with compressed air for 30 seconds. The aeration rate was reduced to bubbling at 60 mL per minute. The counter electrode, salt bridge, and reference electrode were then put in place. The sample substrate was prepared by applying a nonconductive, non-dissolving gel (Nyogel 759G, Nyc Lubricants, Inc.) on the back, edges, and part of the front of the panel to leave a 2 cm x 2 cm test area uncoated for the polarization test. Finally the test sample (working electrode) was placed in the cell and positioned approximately 2-3 mm from the tip of the salt bridge. The cell potential vs. Open circuit was monitored to determine when a steady value was reached (approximately 15-30 minutes) via monitoring by CorrWare for Windows, Version 1.4 software (Scribner Associates, Inc. Charlottesville, VA). After the steady state cell potential was achieved, a potentiodynamic polarization scan was performed by scanning from -0.2 mV to +0.2 mV from the steady state cell potential(Ecorr) vs. Open circuit potential. A scan rate of 10 mV/sec was utilized. The relatively fast scan rate was necessary to complete the anodic portion of the scan before enough zinc had dissolved from the surface to expose the underlying steel base material. The scanning software recorded one data point (current density in A/sq. cm) at each millivolt. Calculations were made utilizing CorrView

5 for Windows version 1.4 software (Scribner Associates, Inc. Charlottesville, VA) which both plotted the data (Potential in volts versus Log Current Density in A/sq. cm) , determined anodic (Ba) and cathodic (Bc) Tafel slope constants, and calculated the corrosion current (Icorr) density (intersection of the anodic and cathodic Tafel lines).

The following 4 samples were evaluated:

10 a). Panel #639 was an electrogalvanized steel panel (ACT E60 EZG 60G 2-side, clean, unpolished, 3" x 6" x 0.030") was rinsed with reagent alcohol and coated with a resin which consisted of:

80.5% by weight NeoRez R9637 (Zeneca Resins)
6.5% by weight N grade sodium Silicate (PQ Corp.)
13.0% by weight water

15 The coating was applied with a standard #12 draw down rod followed by a 15 minute oven bake at 60 C. After allowing to cool for 10 minutes, the coating process was repeated to yield a dry film coating thickness of approximately 0.7 mils. After 570 days the panel was immersed in N-methyl-pyrrolidone for 24 hours to dissolve coating. After light wiping , air drying, and reagent alcohol rinsing, a 2 cm x 7.5 cm section was cut off for potentiodynamic polarization testing.

20 b). Panel #4323 was an electrogalvanized steel panel (ACT E60 EZG 60G 2-side, clean, unpolished, 3" x 6" x 0.030") was rinsed with reagent alcohol and then with naphtha and coated with a gel of the following composition:

70.0 wt % Nygel 759G (Nyje Lubricants, Inc.)
10.0 wt % Sodium Silicate (PQ Corp.)
10.0 wt % Kasil ss-powd Potassium Silicate (PQ Corp.)
10.0 wt % Borogard ZB Zinc Borate (U.S. Borax)

25 The coating was applied with a gate type applicator to a thickness of 0.062 inches. After 107 days, the excess gel was carefully wiped off and the remaining thin film of gel was removed by spray-washing with naphtha with a solvent squirt bottle. The panel was allowed to air dry and then a 2 cm x 7.5 cm section was cut off for potentiodynamic polarization testing.

30 c.) Panel #4977 was an electrogalvanized steel panel (ACT E60 EZG 60G 2-side, clean, unpolished, 3" x 6" x 0.030") was rinsed with reagent alcohol and then wiped with a Kimwipe. A 3" x 4.5" area of the panel was then immersed in 800 mL of a solution of the following

5 composition:
10 wt. % N grade sodium silicate solution (PQ Corp.)
90 wt. % deionized water

The panel was connected as the cathode to a DC power supply (Leader model # 718-20D) and a potential of 6.0 volts was applied. The anode was a panel of the same type as the cathode with the same area immersed in the solution (1:1 anode:cathode surface area), approximately 1.25 inches from the cathode. After 1 hour, the power supply was turned off and the panel was removed from solution. The panel was then air dried while hanging in a vertical position. After 24 days a 2 cm x 7.5 cm section was cut off for potentiodynamic polarization testing.

d.) An unratined an electrogalvanized steel panel (ACT E60 EZG 60G 2-side, clean, unpolished, 3" x 6" x 0.030") was rinsed with reagent alcohol, wiped with a Kimwipe and then rinsed with naphtha and wiped with a Kimwipe and used as a control sample. A 2 cm x 7.5 cm section was cut off for potentiodynamic polarization testing.

The following results were obtained:

Sample #	Ecorr (Volts)	Icorr (A/sq. cm)	Bc	Ba
A	-1.0275	2.27876 x 10 ⁻³	-0.142857	0.021357
B	-1.0491	8.15915 x 10 ⁻³	-0.305985	0.027159
C	-1.0538	1.74087 x 10 ⁻³	-0.583354	0.026380
Control	-1.0408	5.62732 x 10 ⁻³	-0.809446	0.045221

The above results demonstrate that the aforementioned treatments produce a surface with a significantly lower corrosion current density (Icorr) value and thus are significantly more corrosion resistant.

EXAMPLE 36

The following Example demonstrates formation of the previously described mineral layer as a result of a component of the grease/gel interacting with the surface of galvanize metal

5 substrates. The interaction was detected by using ESCA analysis in accordance with conventional methods.

Analytical conditions for ESCA:

10	Instrument X-ray source Source power Analysis region Exit angle*	Physical Electronics Model 5701 LSci Monochromatic aluminum 350 watts 2 mm X 0.8 mm 50° ±7°
15	Electron acceptance angle Charge neutralization Charge correction	electron flood gun C-(C ₂ H) in C 1s spectra at 284.6 eV

• Exit angle is defined as the angle between the sample plane and the electron analyzer lens.

Coatings were made up based on the ingredients and formulation methods shown in Example 10. Different base oils and base oil combinations, alkali silicate types, silicate amounts, and substrates were used to represent a cross section of possible ranges. The different base oils comprised polyalphaolefin (polymerized 1-decene) and linseed oil. Two types of alkali silicates were also used, sodium and calcium silicate. The concentration of the alkali silicate was also varied from 1% to 50% wt to show the range of possible concentrations. Each set of coatings were applied onto both cold rolled and galvanized steel panels.

Each formulation was mixed together and applied onto the given substrate at a thickness between 5 and 10 mils. The coatings were allowed to set for at least 24 hours and then removed from the substrate. Removal was accomplished by first scraping off the excess coating. The residual coating was washed with the base oil used in the formulation to absorb any of the silica or silicates. Finally the excess oil is removed by washing with copious amounts of naphtha. Not adequately removing the silica from the residual coating, will leave behind a precipitate in the subsequent naphtha washing, making any surface analysis more difficult to impossible.

Formulations used for ESCA/XPS analysis

Sample #	1	2	3	4	5	6	7	8
(PAO)								
Durasyn 174	49.3	44.3	49.3	44.3	87	79.2	70.4	44
Linseed Oil	49	44	49	44	0	0	0	0
wt.%								
Fumed Silica	0.7	0.7	0.7	0.7	12	10.8	9.6	6
wt.%								
Sodium silicate	1	10	0	0	0	20	50	
wt.%								

Formulations used for ESCA/XPS analysis on lead panels

Sample #	1	2	3	4
Durasyn 174	89	74	89	44
wt.%				
Fumed Silica	6	6	6	6

wt.%	Sodium silicate	0	0	5	50	wt.%	Fluorine Silica	6	6	6	6
wt.%	Calcium silicate	5	20	0	0	wt.%	Sodium silicate	0	0	5	50
wt.%						wt.%	Calcium silicate	5	20	0	0
wt.%						wt.%					

5 ESCA was used to analyze the surface of each of the substrates. ESCA detects the reaction products between the metal substrate and the coating. Every sample measured showed a mixture of silica and metal silicate. The metal silicate is a result of the reaction between the metal cations of the surface and the alkali silicates of the coating. The silica is a result of either excess silicates from the reaction or precipitated silica from the coating removal process. The metal silicate is indicated by a Si (2p) binding energy (BE) in the low 102 eV range, typically between 102.1 to 102.3. The silica can be seen by Si(2p) BE between 103.3 to 103.6 eV. The resulting spectra show some overlapping peaks, upon deconvolution reveal binding energies in the ranges representative of metal silicate and silica.

10 samples were in the range of 102.1 to 102.3 eV.

EXAMPLE 38

The following Example demonstrates formation of the previously described mineral layer as a result of a component of the grease/gel interacting with the surface of Galfan® substrates (a commercially available alloy comprising zinc and aluminum). The interaction was detected by using ESCA analysis in accordance with conventional methods.

20 Coatings were made up based on the ingredients shown in table shown below. Different alkali silicate types and silicate amounts were used to represent a cross section of possible ranges. Two types of alkali silicates were also used, sodium and calcium silicate. The concentration of the alkali silicate was also varied from 5% to 50% wt to show the range of possible concentrations. Each coatings was applied onto galfan coated steel coupons. Prior to gel application, the copper coupons cut from copper sheets (C110, Fullerton Metals) were rinsed with reagent alcohol.

Each formulation was mixed together and applied onto a lead coupon at a thickness between 5 and 10 mils. The coatings were allowed to set for at least 24 hours and then removed from the substrate. Removal was accomplished by first scraping off the excess coating. The residual coating was washed with the base oil used in the formulation to absorb any of the silica or silicates. Finally the excess oil is removed by washing with copious amounts of naphtha. Not adequately removing the silica from the residual coating, will leave behind a precipitate in the subsequent naphtha washing, making any surface analysis more difficult to impossible.

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Formulations used for ESCA/XPS analysis on Galfan® panels

Sample #	1	2	3	4
Dursyn 174	89	74	89	44

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EXAMPLE 39

The following Example demonstrates formation of the previously described mineral layer as a result of a component of the grease/gel interacting with the surface of copper substrates. The interaction was detected by using ESCA analysis in accordance with conventional methods.

20 Coatings were made up based on the ingredients shown in table shown below. Different alkali silicate types and silicate amounts were used to represent a cross section of possible ranges. Two types of alkali silicates were also used, sodium and calcium silicate. The concentration of the alkali silicate was also varied from 5% to 50% wt to show the range of possible concentrations. Each coatings was applied onto galfan coated steel coupons. Prior to gel application, the copper coupons cut from copper sheets (C110, Fullerton Metals) were rinsed with reagent alcohol.

Each formulation was mixed together and applied onto a lead coupon at a thickness between 5 and 10 mils. The coatings were allowed to set for at least 24 hours and then removed from the substrate. Removal was accomplished by first scraping off the excess coating. The residual coating was washed with the base oil used in the formulation to absorb any of the silica or silicates. Finally the excess oil is removed by washing with copious amounts of naphtha. Not adequately removing the silica from the residual coating, will leave behind a precipitate in the subsequent naphtha washing, making any surface analysis more difficult to impossible.

15 The following Example demonstrates formation of the previously described mineral layer as a result of a component of the grease/gel interacting with the surface of copper substrates. The interaction was detected by using ESCA analysis in accordance with conventional methods.

20 Coatings were made up based on the ingredients shown in table shown below. Different alkali silicate types and silicate amounts were used to represent a cross section of possible ranges. Two types of alkali silicates were also used, sodium and calcium silicate. The concentration of the alkali silicate was also varied from 5% to 50% wt to show the range of possible concentrations. Each coatings was applied onto galfan coated steel coupons. Prior to gel application, the copper coupons cut from copper sheets (C110, Fullerton Metals) were rinsed with reagent alcohol.

Each formulation was mixed together and applied onto a lead coupon at a thickness between 5 and 10 mils. The coatings were allowed to set for at least 24 hours and then removed from the substrate. Removal was accomplished by first scraping off the excess coating. The residual coating was washed with the base oil used in the formulation to absorb any of the silica or silicates. Finally the excess oil is removed by washing with copious amounts of naphtha. Not adequately removing the silica from the residual coating, will leave behind a precipitate in the subsequent naphtha washing, making any surface analysis more difficult to impossible.

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wt.%	Fumed Silica	6	6	6	6
wt.%	Sodium silicate	0	0	5	50
wt.%	Calcium silicate	5	20	0	0
wt.%				$A_x B_y O_z \cdot nH_2O$	

5 ESCA was used to analyze the surface of each of the substrates. ESCA detects the reaction products between the metal substrate and the coating. Every sample measured showed a mixture of silica and metal silicate. The metal silicate is a result of the reaction between the metal cations of the surface and the alkali silicates of the coating. The silica is a result of either excess silicates from the reaction or precipitated silica from the coating removal process. The metal silicate is indicated by a Si (2p) binding energy (BE) in the low 102 eV range, typically between 102.1 to 102.3. The silica can be seen by Si (2p) BE between 103.3 to 103.6 eV. The resulting spectra show some overlapping peaks, upon deconvolution reveal binding energies in the ranges representative of metal silicate and silica. It is to be understood that the foregoing is illustrative only and that other means and techniques may be employed without departing from the spirit or scope of the invention as defined in the following claims.

5 **THE FOLLOWING IS CLAIMED:**

1. An amorphous mineralized surface comprising a cation and an inorganic oxide wherein the amount of oxygen is less than the stoichiometric.
2. A mineralized layer consisting essentially of the oxide network phase

10 wherein A comprises a modifier cation comprising at least one member selected from the group of Group I, II and III metals, B comprises a network forming cation wherein the values of x, y and z are greater than 0. So long as the ratio of y:z is less than or equal to 4:1.

15 3. A transparent corrosion resistant mineralized layer comprising at least one phase obtained by contacting a silicate containing material with a metal surface at a pH ranging from about 9 to at least 11 wherein the amount of oxygen is less than stoichiometric.

15 4. The mineralized layer of Claim 3 wherein the metal surface comprises at least one member selected from the group consisting of iron, steel, zinc, magnesium, aluminum, vanadium, calcium, beryllium, manganese, cobalt, nickel, copper, brass, bronze, zirconium, thallium, chromium, and alloys thereof.

20 5. The mineralized surface of Claim 1 in which cations comprise one or more metals selected from the class consisting Group I, Group II, Group III, transition metals and rare earth metals of the Periodic Chart of the Elements.

20 6. The mineralized surface of Claim 5 in which the cations comprise an alkaline earth metal.

25 7. The mineralized layer of Claim 2 in which anions comprise at least one member selected from the group consisting of selected from the group consisting of one or more of the anions selected from the group consisting of water soluble salts and/or oxides of tungsten, molybdenum, chromium, titanium, zirconium, vanadium, phosphorus, aluminum, iron, boron, bismuth, gallium, tellurium, germanium, antimony, niobium (also known as columbium), 30 magnesium and manganese, mixtures thereof, among others, and more especially, salts and oxides of silicon, aluminum and iron can be employed

30 8. The mineralized layer of Claim 4 wherein the metal surface comprises zinc and the silicate comprises sodium silicate.

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5 9. The mineralized layer of Claim 2 wherein the contacting comprises providing the

silicate to the metal surface in the form of at least one member selected from the group consisting of a paint, coating or gel.

10. The mineralized surface of Claim 1 wherein the surface consist essentially of a complex oxide of the form:

10 $M_xN_yO_z$, wherein "M" represents one or more cationic elements having a covalency factor of less than about 0.5 that functions to balance the charge of the complex oxide, "N" represents one or more lattice forming elements having a covalency factor greater than about 0.15 that functions as the structural component of the complex oxide and optionally wherein the N_yO_z carries single or multiple crystal structures; and wherein x, y and z comprise any number the total of which balances the charge of the complex oxide.

11. The mineralized surface of Claim 10 wherein the covalency factor of M is less than about 0.33 and the covalency factor N is greater than about 0.33.

12. The mineralized surface of Claim 3 wherein the metal surface comprises zinc and the silicate comprises sodium silicate.

13. The mineralized surface of Claim 3 wherein the carrier comprises PAO or polyurethane and the silicate comprises about 1 to 30 wt.% of the carrier.

14. The mineralized surface of Claim 12 wherein the surface is heat treated at a temperature of about 125 to about 175 C.

15. The mineralized surface of Claim 1 wherein the Si(2p) photoelectron binding energy, measured by X-ray Photoelectron Spectroscopy, is higher than 102.1, but less than 103.3 eV, wherein the spectra identifies the mineralized species and wherein the binding energy is higher due to the accumulation of a silica oxide network.

A. CLASSIFICATION OF SUBJECT MATTER		
IPC6: C09D 5/08 According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED		
Minimum documentation searched (classification system followed by classification symbols)		
IPC6: C09D Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)		
WPI		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 9612770 A1 (ELISHA TECHNOLOGIES CO. L.L.C.), 2 May 1996 (02.05.96); page 11, line 18 - page 12, line 2; page 20, line 1 - line 9, claims 1,9,16	1-15
X	US 5068134 A (JAMES A. COLE ET AL.), 26 November 1991 (26.11.91); column 2, line 35 - line 50; column 3, line 59 - line 66, claim 1	1-15
X	US 4842645 A (SHIGEO MIYATA ET AL.), 27 June 1989 (27.06.89), the claims	1-7, 10-11
<input checked="" type="checkbox"/> Further documents are listed in the continuation of Box C.		<input checked="" type="checkbox"/> See parent family annex.
<p>* Special categories of cited documents:</p> <ul style="list-style-type: none"> • Special categories of cited documents • "T" later document published after the international filing date, or priority date and not in conflict with the international application, cited to determine the patentability of the application. • "X" document of particular relevance to the claimed invention, cannot be considered, nor can it be considered to involve an inventive step when the document is taken alone • "Y" document of particular relevance to the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art • "Z" document published prior to the international filing date but later than the priority date claimed • "A" document member of the same patent family 		
Date of the actual completion of the international search		Date of mailing of the international search report
25 May 1998		25.06.1998
		Authorized officer BARBRO NILSSON

Form PCT/ISA210 (second sheet) (July 1992)

2
INTERNATIONAL SEARCH REPORT

International application No.
PCT/US 98/01772

C (Continuation), DOCUMENTS CONSIDERED TO BE RELEVANT	
Category	Classification of document, with indication, where appropriate, of the relevant passages
X	US 4645790 A (GARY T. FREY ET AL), 24 February 1987 (24.02.87), column 2, line 25 - line 34, claims 1-5
X	DE 3618841 A1 (DAIMLER-BENZ AG), 31 March 1988 (31.03.88), column 1, line 37 - line 57, claims 1 and 5

INTERNATIONAL SEARCH REPORT
Information on patent family members

29/04/98

International application No.
PCT/US 98/01772

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US 4842645 A	27/06/89	DE 388691 D, T EP 028953 A, B JP 1891079 C JP 6015679 B JP 63273670 A	19/05/94 09/11/88 07/12/94 02/03/94 10/11/88
US 4645790 A	24/02/87	CA 1240307 A US 4555445 A	09/08/88 26/11/85
DE 3618841 A1	31/03/88	NONE	